

JS011720020B2

(12) United States Patent

Hatakeyama

(10) Patent No.: US 11,720,020 B2

(45) **Date of Patent:**

Aug. 8, 2023

(54) RESIST COMPOSITION AND PATTERNING PROCESS

- (71) Applicant: Shin-Etsu Chemical Co., Ltd., Tokyo
 - (JP)
- (72) Inventor: Jun Hatakeyama, Joetsu (JP)
- (73) Assignee: SHIN-ETSU CHEMICAL CO., LTD.,
 - Tokyo (JP)
- (*) Notice: Subject to any disclaimer, the term of this

patent is extended or adjusted under 35

U.S.C. 154(b) by 440 days.

- (21) Appl. No.: 16/984,535
- (22) Filed: Aug. 4, 2020
- (65) Prior Publication Data

US 2021/0063879 A1 Mar. 4, 2021

(30) Foreign Application Priority Data

Sep. 4, 2019 (JP) 2019-160863

(51) Int. Cl. G03F 7/038 (2006.01) G03F 7/26 (2006.01) G03F 7/00 (2006.01) G03F 7/039 (2006.01)

(52) U.S. Cl.

CPC G03F 7/0382 (2013.01); G03F 7/0002 (2013.01); G03F 7/0392 (2013.01); G03F 7/26 (2013.01)

(58) Field of Classification Search

CPC G03F 7/0382; G03F 7/0002; G03F 7/0392; G03F 7/26; G03F 7/0397; G03F 7/0045; G03F 7/004; G03F 7/033; G03F 7/0395; G03F 7/2004

(56) References Cited

U.S. PATENT DOCUMENTS

4,533,773	A	8/1985	Michaelson et al.
6,294,693	В1	9/2001	Asakawa et al.
7,960,091	B2	6/2011	Shimizu et al.
9,005,874	B2	4/2015	Komuro et al.
9,122,153	B2	9/2015	Echigo et al.
9,176,379	B2	11/2015	Ichikawa et al.
9,250,518	B2	2/2016	Hatakeyama et al.
9,360,753	B2	6/2016	Hatakeyama
9,448,475	B2	9/2016	Masuyama et al.
9,563,123	B2	2/2017	Masuyama et al.
10,101,654	B2	10/2018	Hatakeyama et al.
10,295,904	B2	5/2019	Hatakeyama et al.
10,303,056	B2	5/2019	Hatakeyama et al.
10,474,030	B2	11/2019	Hatakeyama et al.
2003/0152190	A1	8/2003	Watanabe et al.
2003/0198894	$\mathbf{A}1$	10/2003	Mizutani et al.
2007/0043234	A1	2/2007	Vaultier et al.
2007/0259773	A1	11/2007	Burdeniuc et al.
2012/0082936	A1	4/2012	Serizawa et al.
2012/0149916	A1	6/2012	Utsumi et al.

	_ /	
2012/0208127 A1	8/2012	Hatakeyama
2013/0029270 A1	1/2013	Hatakeyama
2013/0052588 A1	2/2013	Yoshida et al.
2014/0242526 A1	8/2014	Allen et al.
2015/0086926 A1	3/2015	Ohashi et al.
2016/0048076 A1	2/2016	Hatakeyama et al.
2017/0174801 A1	6/2017	Hirano
2017/0184962 A1	6/2017	Hatakeyama et al.
2017/0299963 A1	10/2017	Fujiwara
2017/0315442 A1	11/2017	Fukushima et al.
2017/0351177 A1	12/2017	Hatakeyama et al.
2018/0081267 A1	3/2018	Hatakeyama et al.
2018/0081268 A1	3/2018	Hatakeyama
2018/0088463 A1	3/2018	Hatakeyama et al.
2018/0101094 A1	4/2018	Hatakeyama et al.
2018/0143532 A1	5/2018	Hatakeyama
2018/0364570 A1	12/2018	Hatakeyama et al.
2018/0373148 A1	12/2018	Hatakeyama et al.
2019/0155155 A1	5/2019	Hatakeyama et al.
2020/0241414 A1	7/2020	Hatakeyama et al.
2020/0241418 A1	7/2020	Hatakeyama
2020/0272048 A1	8/2020	Hatakeyama et al.
2020/0301274 A1	9/2020	Taniguchi et al.
2021/0033970 A1	2/2021	Hatakeyama et al.
2021/0033971 A1	2/2021	Hatakeyama
2021/0063879 A1	3/2021	Hatakeyama
2021/0149301 A1	5/2021	Fujiwara et al.
2021/0179554 A1	6/2021	Fujiwara et al.
2021/0188770 A1	6/2021	Fujiwara et al.
		3

FOREIGN PATENT DOCUMENTS

JP	H11-102072 A	4/1999
JP	2001-194776 A	7/2001
JP	2002-226470 A	8/2002
	(Cont	inued)

OTHER PUBLICATIONS

Notice of Allowance dated Jun. 29, 2022, issued in U.S. Appl. No. 16/919,574. (13 pages).

Eckert et al., "CIDNP Investigation of Radical Decay Pathways in the Sensitized Photolysis of Triphenylsulfonium Salts", J. Am. Chem. Soc.,1999, 121, 10, pp. 2274-2280. (7 pages).

Wang et al., "Photobase generator and photo decomposable quencher for high-resolution photoresist applications", Proc. of SPIE, 2010, vol. 7639, pp. 76390W1-76390W15. (15 pages).

(Continued)

Primary Examiner — Caleen O Sullivan (74) Attorney, Agent, or Firm — WHDA, LLP

(57) ABSTRACT

A resist composition comprising a base polymer and a salt is provided. The salt consisting of an anion derived from a carboxylic acid having an iodized or brominated hydrocarbyl group and a cation derived from a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound. The resist composition exerts a high sensitizing effect and an acid diffusion suppressing effect, causes no film thickness loss after development, and is improved in resolution, LWR and CDU when a pattern is formed therefrom by lithography.

13 Claims, No Drawings

US 11,720,020 B2 Page 2

(56)		ces Cited	KR KR TW	10-2012-0093777 A 10-2016-0019860 A 201241556 A	8/2012 2/2016 10/2012
	FOREIGN PALE	NT DOCUMENTS	TW	201516024 A	5/2015
-	2002 252440	12/2022	TW	201730674 A	9/2017
JP	2002-363148 A	12/2002	WO	2008/066011 A1	6/2008
JP	2008-133312 A	6/2008	WO	2010/059174 A1	5/2010
JР	2009-145578 A	7/2009	WO	2013/024777 A1	2/2013
JР	2009-181062 A	8/2009			
JР	2010-061087 A	3/2010		OTHER BILL	OLIC ATIONS
JР	2011-39266 A	2/2011			BLICATIONS
JР	2011-039502 A	2/2011		,	1, issued in TW Application No.
JР	2011-530652 A	12/2011	109127	7275 (counterpart to U.S. A	Appl. No. 16/871,648). (10 pages).
JP	2013-025211 A	2/2013	Non-Fi	inal Office Action dated M	Iar. 29, 2022, issued in U.S. Appl.
JР	2013-83957 A	5/2013	No. 16	/871,648. (23 pages).	**
JР	2015-90382 A	5/2015	Office	Action dated Oct. 15, 2	2018, issued in TW Application
JР	2015-161823 A	9/2015			Appl. No. 15/725,404). (9 pages).
JР	2015-172746 A	10/2015			8, issued in KR Application No.
JР	2015-180928 A	10/2015			U.S. Appl. No. 15/725,404), with
JP	5852490 B2	2/2016			U.S. Appl. No. 13/723,404), with
JР	2017-219836 A	12/2017		translation. (13 pages).	
JР	2018-4812 A	1/2018			Iar. 22, 2019, issued in U.S. Appl.
JP	2018-49264 A	3/2018		/725,404. (28 pages).	
JР	2018-060069 A	4/2018	Office	Action dated Apr. 16, 2	2021, issued in counterpart TW
JP	2018-97356 A	6/2018	Applic	ation No. 109129970. (8 p	pages).

RESIST COMPOSITION AND PATTERNING PROCESS

CROSS-REFERENCE TO RELATED APPLICATION

This non-provisional application claims priority under 35 U.S.C. § 119(a) on Patent Application No. 2019-160863 filed in Japan on Sep. 4, 2019, the entire contents of which are hereby incorporated by reference.

TECHNICAL FIELD

This invention relates to a resist composition and a pattern forming process.

BACKGROUND ART

To meet the demand for higher integration density and operating speed of LSIs, the effort to reduce the pattern rule 20 is in rapid progress. In particular, the enlargement of the logic memory market to comply with the wide-spread use of smartphones drives forward the miniaturization technology. As the advanced miniaturization technology, manufacturing of microelectronic devices at the 10-nm node by double 25 patterning of the ArF immersion lithography has been implemented in a mass scale. Manufacturing of 7-nm node devices as the next generation by the double patterning technology is approaching to the verge of high-volume application. The candidate for 5-nm node devices as the next generation but one is EUV lithography.

As the pattern feature size is reduced, approaching to the diffraction limit of light, light contrast lowers. In the case of positive resist film, a lowering of light contrast leads to reductions of resolution and focus margin of hole and trench 35 patterns. For preventing the reduction of resolution of resist pattern due to a lowering of light contrast, an attempt is made to enhance the dissolution contrast of resist film.

Chemically amplified resist compositions comprising an acid generator capable of generating an acid upon exposure 40 to light or EB include chemically amplified positive resist compositions wherein deprotection reaction takes place under the action of acid and chemically amplified negative resist compositions wherein polarity switch or crosslinking reaction takes place under the action of acid. Quenchers are 45 often added to these resist compositions for the purpose of controlling the diffusion of the acid to unexposed region to improve the contrast. The addition of quenchers is fully effective to this purpose. A number of amine quenchers were proposed as disclosed in Patent Documents 1 to 3.

With respect to the acid labile group used in (meth) acrylate polymers for the ArF lithography resist material, deprotection reaction takes place when a photoacid generator capable of generating a sulfonic acid having fluorine substituted at α-position (referred to "α-fluorinated sulfonic 55 acid") is used, but not when an acid generator capable of generating to a sulfonic acid not having fluorine substituted at α -position (referred to " α -non-fluorinated sulfonic acid") or carboxylic acid is used. If a sulfonium or iodonium salt capable of generating an α -fluorinated sulfonic acid is 60combined with a sulfonium or iodonium salt capable of generating an α-non-fluorinated sulfonic acid, the sulfonium or iodonium salt capable of generating an α-non-fluorinated sulfonic acid undergoes ion exchange with the α -fluorinated sulfonic acid. Through the ion exchange, the α-fluorinated 65 sulfonic acid thus generated by light exposure is converted back to the sulfonium or iodonium salt while the sulfonium

2

or iodonium salt of an α -non-fluorinated sulfonic acid or carboxylic acid functions as a quencher. Patent Document 4 discloses a resist composition comprising a sulfonium or iodonium salt capable of generating carboxylic acid as a quencher.

Patent Document 5 discloses a resist composition comprising a sulfonium salt of iodized aromatic carboxylic acid as a quencher. The quencher is readily decomposed upon EUV exposure due to the substantial EUV absorption of iodine, and is quite effective for controlling acid diffusion by virtue of the large atomic weight of iodine. Thus a high sensitivity and an improved dimension uniformity due to low acid diffusion are expectable.

Patent Document 6 discloses a resist composition comprising an iodized aniline as a quencher. The aniline has a low basicity which is insufficient to suppress acid diffusion.

Sulfonium and iodonium salt type quenchers are photodecomposable like photoacid generators. That is, the amount of quencher in the exposed region is reduced. Since acid is generated in the exposed region, the reduced amount of quencher leads to a relatively increased concentration of acid and hence, an improved contrast. However, the acid diffusion in the exposed region is not suppressed, indicating the difficulty of acid diffusion control.

Patent Document 7 discloses a resist composition comprising a 2,5,8,9-tetraaza-1-phophabicyclo[3.3.3]undecane salt, biguanide salt, and phosphazene salt of an iodized aromatic carboxylic acid. A sensitizing effect due to the high absorption of iodine is expectable as well as a high sensitivity and low acid diffusion assigned to a salt of a strong base or 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound. Nevertheless, there is the demand for a resist material of new concept which exhibits a higher sensitivity and lower acid diffusion.

CITATION LIST

Patent Document 1: JP-A 2001-194776
Patent Document 2: JP-A 2002-226470
Patent Document 3: JP-A 2002-363148

Patent Document 4: WO 2008/066011 Patent Document 5: JP-A 2017-219836

Patent Document 6: JP-A 2018-097356 Patent Document 7: JP-A 2018-049264 (U.S. Pat. No. 10,101,654)

SUMMARY OF INVENTION

As the wavelength of light becomes shorter, the energy density thereof becomes higher and hence, the number of photons generated upon exposure becomes smaller. A variation in photon number causes variations in the LWR of line patterns and the CDU of hole patterns. As the exposure dose increases, the number of photons increases, leading to a less variation of photon number. Tis there is a tradeoff relationship between sensitivity and resolution, LWR and CDU. In particular, the EUV lithography resist materials have the tendency that a lower sensitivity leads to better LWR and CDU.

An increase in acid diffusion also causes degradation of resolution, LWR and CDU. This is because acid diffusion not only causes image blur, but also proceeds non-uniformly in a resist film. For suppressing acid diffusion, it is effective to lower the PEB temperature, to use a bulky acid which is least diffusive, or to increase the amount of quencher added. However, any of these means for reducing acid diffusion results in a lowering of sensitivity. Either the means for

reducing photon variation or the means for reducing acid diffusion variation leads to a lowering of resist sensitivity.

An object of the invention is to provide a resist composition which has a high sensitizing effect and an acid diffusion suppressing effect and causes no film thickness loss after development and which has improved resolution, LWR and CDU, and a pattern forming process using the same.

A significant increase of acid generation efficiency and a significant suppression of acid diffusion must be achieved before the tradeoff relationship between sensitivity and resolution, LWR and CDU can be overcome.

The inventor has found that when a salt consisting of an anion derived from a carboxylic acid having an iodized or brominated hydrocarbyl group exclusive of iodized or brominated aromatic ring and a cation derived from a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound is added to a base polymer, the resulting resist composition forms a resist film which exerts a high sensitizing effect and an acid diffusion suppressing effect and has a high sensitivity, minimized LWR and improved CDU.

In one aspect, the invention provides a resist composition comprising a base polymer and a salt, the salt consisting of an anion derived from a carboxylic acid having an iodized or brominated hydrocarbyl group exclusive of iodized or brominated aromatic ring and a cation derived from a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound.

Typically, the salt has the formula (A).

$$\left[\left(X^{RI}\right)_{n}X^{1}\right]_{m}X^{2} \xrightarrow{O^{-}} A^{+}$$

Herein m and n are each independently 1, 2 or 3. X^{BI} is iodine or bromine. X^1 is a single bond, ether bond, ester bond, amide bond, carbonyl group or carbonate group. X^2 is a single bond or a C_1 - C_{20} (m+1)-valent hydrocarbon group which may contain a heteroatom other than iodine and bromine. R^1 is a C_1 - C_{20} (n+1)-valent aliphatic hydrocarbon group which may contain at least one moiety selected from the group consisting of fluorine, chlorine, hydroxyl, carboxyl, C_6 - C_{12} aryl, ether bond, ester bond, carbonyl, amide bond, carbonate, urethane bond, and urea bond. A^+ is a cation having the formula (A)-1, (A)-2 or (A)-3.

4

-continued

Herein R¹¹ to R¹³ are each independently a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom. R¹⁴ to R²¹ are each independently hydrogen or a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom, a pair of R¹⁴ and R¹⁵, R¹⁵ and R¹⁶, R¹⁶ and R¹⁷, R¹⁷ and R¹⁸, R¹⁸ and R¹⁹, R¹⁹ and R²⁰, or R²⁰ and R²¹ may bond together to form a ring with the nitrogen atom to which they are attached, or the nitrogen atoms to which they are attached and intervening carbon atom, the ring optionally containing an ether bond. R²² to R²⁹ are each independently hydrogen or a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom, a pair of R²² and R²³, R²³ and R²⁴, R²⁴ and R²⁵, R²⁵ and R²⁶, R²⁶ and R²⁷, or R²⁷ and R²⁸ may bond together to form a ring with the nitrogen atom to which they are attached and intervening phosphorus atom, a pair of R²² and R²³, R²⁴ and R²⁵, R²⁶ and R²⁷, or R²⁸ and R²⁹ may bond together to form a group having the formula (A)-3-1, and R² may be a group having the formula (A)-3-2 when R²² is hydrogen.

$$R^{36}$$
 R^{37}
 $P = 0$
 R^{39}
 R^{38}
 R^{38}

(A)-1 Herein R³⁰ to R³⁹ are each independently hydrogen or a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom, a pair of R³⁰ and R³¹, R³¹ and R³², R³² and R³³, R³³ and R³⁴, R³⁴ and R³⁵, R³⁶ and R³⁷, or R³⁸ and R³⁹ may bond together to form a ring with the nitrogen atom to which they are attached, or the nitrogen atoms to which they are attached and intervening phosphorus atom, a pair of R³⁰ and R³¹, R³² and R³³, or R³⁴ and R³⁵ may bond together to form a group having the formula (A)-3-I. The broken line designates a valence bond.

The resist composition may further comprise an acid generator capable of generating a sulfonic acid, sulfonimide or sulfonemethide.

In a preferred embodiment, the base polymer comprises recurring units having the formula (a1) or recurring units having the formula (a2).

Herein R^4 is each independently hydrogen or methyl. R^{41} and R^{42} each are an acid labile group, Y^1 is a single bond, phenylene group, naphthylene group, or C_1 - C_{12} linking $_{25}$ group containing at least one moiety selected from ester bond and lactone ring, and Y^2 is a single bond or ester bond.

Typically the resist composition is a chemically amplified positive resist composition.

In another embodiment, the base polymer is free of an 30 acid labile group. Typically the resist composition is a chemically amplified negative resist composition.

In a further preferred embodiment, the base polymer comprises recurring units of at least one type selected from recurring units having the formulae (f1) to (f3).

$$\begin{array}{ccc}
F_2 & R^{55} \\
& R^{55}
\end{array}$$

$$\begin{array}{cccc}
& R^{56} \\
& SO_3 & SO_3 & R^{56} \\
& R^{57} & R^{57} \\
& R^{58} & R^{57}
\end{array}$$

Herein R^A is each independently hydrogen or methyl. Z^1 is 65 a single bond, phenylene group, $-O-Z^{11}-$, -C(=O)- $O-Z^{11}-$ or $-C(=O)-NH-Z^{11}-$, Z^{11} is a C_1 - C_6 ali-

phatic hydrocarbylene group or phenylene group, which may contain a carbonyl, ester bond, ether bond or hydroxyl moiety. Z² is a single bond, —Z²¹—C(=O)—O—, —Z²¹—O— or —Z²¹—O—C(=O)—, Z²¹ is a C₁-C₁₂ saturated hydrocarbylene group which may contain a carbonyl moiety, ester bond or ether bond. Z³ is a single bond, methylene, ethylene, phenylene, fluorinated phenylene, —O—Z³¹—, —C(=O)—O—Z³¹—, or —C(=O)—NH—Z³¹—, Z³¹ is a C₁-C₆ aliphatic hydrocarbylene group, phenylene group, fluorinated phenylene group, or trifluoromethyl-substituted phenylene group, which may contain a carbonyl moiety, ester bond, ether bond or hydroxyl moiety. R⁵¹ to R⁵² are each independently a C₁-C₂o hydrocarbyl group which may contain a heteroatom, any two of R⁵³, R⁵³ and R⁵⁵ or any two of R⁵⁵6, R⁵⁵7 and R⁵⁵8 may bond together to form a ring with the sulfur atom to which they are attached. R^{HF} is hydrogen or trifluoromethyl. M⁻ is a non-nucleophilic counter ion.

The resist composition may further comprise an organic solvent and/or a surfactant.

In another aspect, the invention provides a process for forming a pattern comprising the steps of applying the resist composition defined above to form a resist film on a substrate, exposing the resist film to high-energy radiation, and developing the exposed resist film in a developer.

Preferably, the high-energy radiation is ArF excimer laser radiation of wavelength 193 n, KrF excimer laser radiation of wavelength 248 nm, EB or EUV of wavelength 3 to 15 nm

Advantageous Effects of Invention

A resist film contains the salt of a carboxylic acid having an iodized or brominated hydrocarbyl group (exclusive of 35 iodized or brominated aromatic ring) with a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound. Because of inclusion of iodine or bromine featuring substantial light absorption, the salt exhibits a sensitizing effect due to secondary electrons released therefrom upon exposure. Bulky and strongly basic 2,5,8,9tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide and phosphazene compounds exert an acid diffusion suppressing effect and a high dissolution contrast. Thus the resist film containing the salt exhibits a high resolution, wide focus (f2) 45 margin, high sensitivity, and minimal LWR or improved CDU as a positive or negative resist film subject to aqueous alkaline development or as a negative resist film subject to organic solvent development.

DESCRIPTION OF EMBODIMENTS

As used herein, the singular forms "a," "an" and "the" include plural referents unless the context clearly dictates otherwise. The notation $(C_n - C_m)$ means a group containing from n to m carbon atoms per group. In chemical formulae, the broken line designates a valence bond, Me stands for methyl, Ac for acetyl, and Ph for phenyl. As used herein, the term "iodized" or "brominated" indicates that a compound is substituted with iodine or bromine or a compound contains iodine or bromine.

The abbreviations and acronyms have the following meaning.

EB: electron beam

50

EUV: extreme ultraviolet

Mw: weight average molecular weight Mn: number average molecular weight

Mw/Mn: molecular weight dispersity

GPC: gel permeation chromatography

PEB: post-exposure bake PAG: photoacid generator LWR: line width roughness CDU: critical dimension uniformity

Resist Composition

The resist composition of the invention is defined as comprising a base polymer and a salt, the salt consisting of an anion derived from a carboxylic acid having an iodized or brominated hydrocarbyl group exclusive of iodized or brominated aromatic ring and a cation derived from a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound. The salt is collectively referred to as "iodized or brominated hydrocarbyl-containing carboxylic salt," hereinafter.

The iodized or brominated hydrocarbyl-containing carboxylic salt undergoes ion exchange with sulfonic acid, sulfonimide or sulfonmethide generated from an acid generator, especially sulfonic acid containing fluorinated alky bissulfonimide or trissulfonmethide, whereupon a 2,5,8,9-20 tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene cation form a salt with a fluorinated alkylcontaining sulfonic acid, bissulfonimide or trissulfonmethide and an iodized or brominated hydrocarbyl-containing carboxylic acid is released. The 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene has an acid trapping ability and an acid diffusion suppressing effect. That is, the iodized or brominated hydrocarbylcontaining carboxylic salt functions as a quencher in the resist composition. Since the iodized or brominated hydrocarbyl-containing carboxylic salt is not photosensitive and thus not photo-decomposable, it retains a sufficient acid trapping ability even in the exposed region, suppressing acid diffusion from the exposed region to the unexposed region.

Besides the iodized or brominated hydrocarbyl-containing carboxylic salt, an amine compound, ammonium salt, sulfonium salt or iodonium salt may be separately added as another quencher to the resist composition of the invention. The ammonium salt, sulfonium salt or iodonium salt added as the quencher is preferably an ammonium, sulfonium or iodonium salt of carboxylic acid, sulfonic acid, sulfonamide or saccharin. The carboxylic acid may or may not be fluorinated at α -position.

The acid diffusion suppressing effect and contrast enhancing effect of the iodized or brominated hydrocarbyl-containing carboxylic salt are valid in both the positive or negative pattern formation by aqueous alkaline development and the negative pattern formation by organic solvent development. Iodized or Brominated Hydrocarbyl-Containing Carboxylic Salt

The iodized or brominated hydrocarbyl-containing carboxylic salt typically has the formula (A).

$$\left[\left(X^{BI}\right)_{m}R^{1}-X^{1}\right]_{m}X^{2} \xrightarrow{O^{-}} A^{+}$$

In formula (A), m and n are each independently 1, 2 or 3. X^{BI} is iodine or bromine.

 X^1 is a single bond, ether bond, ester bond, amide bond, carbonyl group or carbonate group.

 $\rm X^2$ is a single bond or a $\rm C_1$ - $\rm C_{20}$ (m+1)-valent hydrocarbon 65 group which may contain a heteroatom other than iodine and bromine.

 R^1 is a C_1 - C_{20} (m+1)-valent aliphatic hydrocarbon group. The aliphatic hydrocarbon group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include alkanediyl groups such as methanediyl, ethane-1,1-diyl, ethane-1,2-diyl, propane-1,1-diyl, propane-1,2-diyl, propane-1,3-diyl, propane-2,2-diyl, butane-1-diyl, butane-1,2-diyl, butane-1,3-diyl, butane-2,3-diyl, butane-1, 4-diyl, 1,1-dimethylethane-1,2-diyl, pentane-1,5-diyl, 2-methylbutane-1,2-diyl, hexane-1,6-diyl, heptane-1,7-diyl, octane-1,8-diyl, nonane-1,9-diyl, decane-1,10-diyl, undecane-1,11-diyl, and dodecane-1,12-diyl; cycloalkanediyl groups such as cyclopropane-1,1-diyl, cyclopropane-1,2diyl, cyclobutane-1,1-diyl, cyclobutane-1,2-diyl, cyclobutane-1,3-diyl, cyclopentane-1,1-diyl, cyclopentane-1,2-diyl, cyclopentane-1,3-diyl, cyclohexane-1,1-diyl, cyclohexane-1,2-diyl, cyclohexane-1,3-diyl, and cyclohexane-1,4-diyl; divalent polycyclic saturated hydrocarbon groups such as norbornane-2,3-diyl and norbomane-2,6-diyl; alkenediyl groups such as 2-propene-1,1-diyl; alkynediyl groups such as 2-propyne-1,1-diyl; cycloalkenediyl groups such as 2-cyclohexene-1,2-diyl, 2-cyclohexene-1,3-diyl, 3-cyclohexene-1,2-diyl; divalent polycyclic unsaturated hydrocarbon groups such as 5-norbornene-2,3-diyl; and cyclic aliphatic hydrocarbon-substituted alkanediyl groups such as cyclopentyhnethanediyl, cyclohexyhnethanediyl, 2-cyclopentenylmethanediyl, 3-cyclopentenylmethanediyl, 2-cyclohexenylmethanediyl, 3-cyclohexenylmethaediyl, as well as trior tetravalent groups obtained by removing one or two hydrogen atoms from the foregoing groups.

In the foregoing groups, some or all of the hydrogen atoms may be substituted by fluorine, chlorine, hydroxyl moiety, carboxyl moiety, or C_6 - C_{12} aryl moiety, and at least one moiety selected from ether bond, ester bond, carbonyl moiety, amide bond, carbonate moiety, urethane bond and urea bond may intervene in a carbon-carbon bond. Suitable C_6 - C_{12} aryl moieties include phenyl, 2-methylphenyl, 3-methylphenyl, 4-methylphenyl, 1-naphthyl, 2-naphthyl and fluorenyl.

Examples of the anion of the salt having formula (A) are given below, but not limited thereto.

-continued

5

10

10

15

20

20

30

35

1

40

45

$$F_2C$$
 F_2C
 F_2C

In formula (A), A^+ is a cation having the formula (A)-1, (A)-2 or (A)-3.

$$\begin{array}{c}
R^{11} \\
N \\
N \\
N
\end{array}$$

$$\begin{array}{c}
H^+ \\
N \\
N
\end{array}$$

$$\begin{array}{c}
R^{12} \\
N \\
N
\end{array}$$

$$\begin{array}{c}
R^{13} \\
\end{array}$$

-continued

In formula (A)-1, R¹⁴ to R²¹ are each independently a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom. In formula (A)-2, R¹⁴ to R²¹ are each independently hydrogen or a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom. A pair of R¹⁴ and R¹⁵, R¹⁵ and R¹⁶, R¹⁶ and R¹⁷, R¹⁷ and R¹⁸, R¹⁸ and R¹⁹, R¹⁹ and R²⁰, or R²⁰ and R²¹ may bond together to form a ring with the nitrogen atom to which they are attached and the intervening carbon atom(s). The ring may contain an ether bond.

In formula (A)-3, R^{22} to R^{29} are each independently hydrogen or a C_1 - C_{24} hydrocarbyl group which may contain a heteroatom. A pair of R^{22} and R^{23} , R^{23} and R^{24} , R^{24} and R^{25} , R^{25} and R^{26} , R^{26} and R^{27} , or R^{27} and R^{28} may bond together to form a ring with the nitrogen atom to which they are attached or the nitrogen atoms to which they are attached and the intervening phosphorus atom. R^{22} and R^{23} , R^{24} and R^{25} , R^{26} and R^{27} , or R^{28} and R^{29} , taken together, may forma group having the formula (A)-3-1. R^{23} may be a group having the formula (A)-3-2 when R^{22} is hydrogen.

In formulae (A)-3-1 and (A)-3-2, R^{30} to R^{39} are each independently hydrogen or a C_1 - C_{24} hydrocarbyl group 50 which may contain a heteroatom. A pair of R^{30} and R^{31} , R^{31} and R^{32} , R^{32} and R^{33} , R^{33} and R^{34} , R^{34} and R^{35} , R^{36} and R^{37} , or R^{38} and R^{39} may bond together to form a ring with the nitrogen atom to which they are attached or the nitrogen atoms to which they are attached and the intervening phosphorus atom. R^{30} and R^{31} , R^{32} and R^{33} , or R^{34} and R^{35} , taken together, may form a group having the formula (A)-3-1.

The C₁-C₂₄ hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include alkyl groups such as methyl, ethyl, n-propyl, 60 isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl n-hexyl, n-heptyl, n-octyl, n-nonyl, n-decyl, n-undecyl and n-dodecyl; cyclic saturated hydrocarbyl groups such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclopentylmethyl, cyclopentylethyl, cyclohexylmethyl, cyclohexylethyl, norbornyl, and adanntyl; alkenyl groups such as vinyl, 1-propenyl, 2-propenyl, 1-butenyl, 2-butenyl, 3-bute-

nyl, pentenyl and hexenyl; alkynyl groups such as ethynyl, 1-propynyl, 2-propynyl, butynyl, pentynyl, and hexynyl; cyclic unsaturated aliphatic hydrocarbyl groups such as cyclopentenyl and cyclohexenyl: aryl groups such as phenyl, methylphenyl ethylphenyl, n-propylphenyl, isopropylphenyl, n-butylphenyl, isobutylphenyl, sec-butylphenyl, tertbutylphenyl, naphthyl, methylnaphthyl, ethylnaphthyl, n-propyInaphthyl, isopropylnaphthyl, n-butynaphthyl, isobutynaphthyl, sec-butylnaphthyl, tert-butylnaphthyl, and fluorenyl; and aralkyl groups such as benzyl, phenethyl, naphthylmethyl, and fluorenylmethyl. In the foregoing groups, some or all hydrogen atoms may be substituted by a moiety containing a heteroatom such as oxygen, sulfur, nitrogen or halogen, and some carbon may be replaced by a moiety containing a heteroatom such as oxygen, sulfur or nitrogen, so that the group may contain a halogen, sulfone, amino, hydroxyl, thiol, nitro, ester bond, ether bond, sulfide bond, sulfoxide, carbonate, carbamate or amide bond.

Examples of the 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3] undecane cation having formula (A)-1 are shown below, but not limited thereto.

Examples of the biguanide cation having formula (A)-2 are shown below, but not limited thereto. $\,$

Examples of the phosphazene cation having formula 35 (A)-3 are shown below, but not limited thereto.

15

20

25

In the cationic 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3] undecane, biguanide or phosphazene compound, positive charges are delocalized among plural nitrogen atoms. Therefore, points of trapping the anion of sulfonic acid, sulfonimide or sulfonmethide for neutralization are distributed everywhere. Thus the anion is quickly trapped. The cationic 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound is an effective quencher having a high basicity and a high trapping ability.

With respect to the synthesis method, the iodized or brominated hydrocarbyl-containing carboxylic salt may be synthesized, for example, by mixing a 2,5,8,9-tetraaza-1-phosphabicyclo[3.3.3]undecane, biguanide or phosphazene compound with an iodized or brominated hydrocarbyl-containing carboxylic acid.

Since the iodized or brominated hydrocarbyl-containing carboxylic salt contains iodine or bromine of relatively large atomic weight in the molecule, it is substantially absorptive to EUV or EB. Iodine or bromine has many electron orbits in its molecule and releases many secondary electrons upon EUV exposure. The secondary electrons thus released provide energy transfer to an acid generator, achieving a high sensitizing effect. This leads to a high sensitivity and low acid diffusion, achieving improvements in both factors of LWR or CDU and sensitivity.

In view of sensitivity and acid diffusion suppressing 55 effect, the iodized or brominated hydrocarbyl-containing carboxylic salt is preferably present in the resist composition in an amount of 0.001 to 50 parts, more preferably 0.01 to 20 parts by weight per 100 pats by weight of the base polymer to be described below.

Base Polymer

In the case of a positive resist composition, the base polymer in the resist composition is a polymer comprising acid labile group-containing recurring units. The acid labile group-containing recurring units are preferably recurring units having the formula (a1) or recurring units having the formula (a2). Sometimes these recurring units are simply referred to as recurring units (a1) and (a2).

$$\begin{array}{c}
 & \text{(al)} \\
 & \text{(al)}$$

$$\begin{array}{c}
 & \text{(a2)} \\
 & \text{V}^{1} \\
 & \text{OR}^{42}
\end{array}$$

Herein R^4 is each independently hydrogen or methyl. R^{41} and R^{42} are each independently an acid labile group. Y^1 is a single bond, phenylene, naphthylene, or a C_1 - C_{12} linking $_{25}$ group containing at least one of ester bond and lactone ring. Y^2 is a single bond or ester bond. R^{41} and R^{42} may be the same or different when the base polymer contains both recurring units (a1) and (a2).

Examples of the monomer from which recurring units 30 (a1) are derived are shown below, but not limited thereto. Herein R^4 and R^{41} are as defined above.

-continued
$$\mathbb{R}^{d}$$
 \mathbb{R}^{d}
 \mathbb{R}^{d}

Examples of the monomer from which recurring units 40 (a2) are derived are shown below, but not limited thereto. Herein R^4 and R^{42} are as defined above.

The acid labile groups represented by R⁴¹ and R⁴² in formulae (a1) and (a2) may be selected from a variety of such groups, for example, those groups described in JP-A

2013-080033 (U.S. Pat. No. 8,574,817) and JP-A 2013-083821 (U.S. Pat. No. 8,846,303).

Typical of the acid labile group are groups of the following formulae (AL-1) to (AL-3).

$$\begin{array}{c}
O \\
\parallel \\
----(CH_2)_a - C - O - \mathbb{R}^{L_1}
\end{array}$$
(AL-1)

$$(AL-3)$$

$$---C$$

$$\downarrow R^{L7}$$

$$\downarrow R^{L6}$$

$$20$$

In formulae (AL-1) and (AL-2), R^{L1} and R^{L2} are each independently a C_1 - C_{40} hydrocarbyl group which may contain a heteroatom such as oxygen, sulfur, nitrogen or fluorine. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Preferred are C_1 - C_{40} , especially C_1 - C_{20} saturated hydrocarbyl groups.

In formula (AL-1), "a" is an integer of 0 to 10, preferably $\ _{30}$ 1 to 5.

In formula (AL-2), R^{L3} and R^{L4} are each independently hydrogen or a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom such as oxygen, sulfur, nitrogen or fluorine. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Preferred are C_1 - C_{24} saturated hydrocarbyl groups. Any two of R^{L2} , R^{L3} and R^{L4} may bond together to form a ring, typically alicyclic, with the carbon atom or carbon and oxygen atoms to which they are attached, the ring containing 3 to 20 carbon atoms, preferably 4 to 16 carbon atoms.

In formula (AL-3), R^{L5} , R^{L6} and R^{L7} are each independently a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom such as oxygen, sulfur, nitrogen or fluorine. The 45 hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Preferred are C_1 - C_{20} saturated hydrocarbyl groups. Any two of R^{L5} , R^{L6} and R^{L7} may bond together to form a ring, typically alicyclic, with the carbon atom to which they are attached, the ring containing 3 to 20 carbon atoms, preferably 4 to 16 carbon atoms.

The base polymer may further comprise recurring units (b) having a phenolic hydroxyl group as an adhesive group. Examples of suitable monomers from which recurring units (b) are derived are given below, but not limited thereto. Herein R⁴ is as defined above.

-continued OH

Further, recurring units (c) having another adhesive group selected from hydroxyl (other than the foregoing phenolic hydroxyl), lactone ring, sultone ring, ether bond, ester bond, sulfonate bond, cabonyl, sulfonyl, cyano and caboxyl groups may also be incorporated in the base polymer. Examples of suitable monomers from which recurring units (c) are derived are given below, but not limited thereto. Herein R^A is as defined above.

$$F_{3}C \longrightarrow CF_{3}$$

$$CF_{3}$$

$$C$$

-continued

$$R^{A}$$
 $F_{3}C$
 $F_{$

In another preferred embodiment, the base polymer may further comprise recurring units (d) derived from indene, benzofuran, benzothiopbene, acenaphthylene, chromone, coumarin, and norbornadiene, or derivatives thereof. Examples of suitable monomers from which reaming units (d) are derived are given below, but not limited thereto.

The base polymer may further include recurring units (e) which are derived from styrene, vinylnaphthalene, vinylanthracene, vinylpyrene, methyleneindene, vinylpyridine, or vinylcarbazole.

In a further embodiment, recurring units (f) derived from an onium salt having a polymerizable unsaturated bond may be incorporated in the base polymer. The preferred recurring units (f) include recurring units having formula (f1), recurring units having formula (f2), and recurring units having formula (f3). These units are simply referred to as recurring units (f1), (2) and (f3), which may be used alone or in combination of two or more types.

(f1)

(f3)

25

30

$$\begin{array}{c}
R^4 \\
\downarrow \\
Z^1 \\
\downarrow \\
R^{51} - S^+ - R^{52} \\
M^-
\end{array}$$

$$Z^{3}$$

$$SO_{3}$$

$$R^{56}$$

$$R^{57}$$

$$R^{58}$$

In formulae (f1) to (f3), R^A is each independently hydrogen or methyl. Z^1 is a single bond, phenylene, $-O-Z^{11}$, $-C(=O)-O-Z^{11}-$, or $-C(=O)-NH-Z^{11}-$, wherein Z^{11} is a C_1 - C_6 aliphatic hydrocarbylene group or 35 phenylene group, which may contain a carbonyl moiety, ester bond, ether bond or hydroxyl moiety. Z2 is a single bond. $-Z^2$ —C(=O)—O—, $-Z^{21}$ —O—, or $-Z^{21}$ —O— C(=O)—, wherein Z^{21} is a C_1 - C_{12} saturated hydrocarbylene group which may contain a carbonyl moiety, ester bond or ether bond. Z³ is a single bond, methylene, ethylene, phenylene, fluorinated phenylene, —O— Z^{31} —, —C(=O)—O— Z^{31} —, or —C(=O)NH— Z^{31} —, wherein Z^{31} is a C_1 - C_6 aliphatic hydrocarbylene group, phenylene, fluorinated phenylene or trifluoromethyl-substituted phenylene group, which may contain a carbonyl moiety, ester bond, ether bond or hydroxyl moiety. The aliphatic hydrocarbylene groups may be saturated or unsaturated and straight, branched or cyclic. The saturated hydrocarbylene groups may be 50 straight, branched or cyclic.

In formulae (f1) to (f3), R^{51} to R^{58} are each independently a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include C_1 - C_{20} alkyl groups C_6 - C_{20} aryl groups, and C_7 - C_{20} aralkyl groups. In these groups, some or all hydrogen atoms may be substituted by C_1 - C_{10} saturated hydrocarbyl, halogen, trifluoromethyl, cyano, nitro, hydroxyl, mercapto, C_1 - C_{10} saturated hydrocarbyloxycarbonyl or C_2 - C_{10} saturated hydrocarbyloxy, or some carbon may be replaced by a carbonyl moiety, ether bond or ester bond. Any two of R^{53} , R^{54} and R^{55} , or any two of R^{56} , R^{57} and R^{58} may bond together to form a ring with the sulfur atom to which they are attached. Examples of the ring are as will be exemplified for the ring that R^{101} and R^{102} in formula

(1-1), taken together, form with the sulfur atom to which they are attached.

In formula (f2), R^{HF} is hydrogen or trifluoromethyl.

In formula (f1), M⁻ is a non-nucleophilic counter ion. Examples thereof include halide ions such as chloride and bromide ions; fluoroalkylsulfonate ions such as triflate, 1,1,1-trifluoroethanesulfonate, and nonafluorobutane-sulfonate; arylsulfonate ions such as tosylate, benzene-sulfonate, 4-fluorobenzenesulfonate, and 1,2,3,4,5-pentafluorobenzenesulfonate; alkylsulfonate ions such as mesylate and butanesulfonate; sulfonimide ions such as bis(trifluoromethylsulfonyl)imide, bis(perfluoroethylsulfonyl)imide and bis(perfluorobutylsulfonyl)imide; and sulfonemethide ions such as tris(trifluoromethylsulfonyl) methide and tris(perfluoroethylsulfonyl)methide.

Also included are sulfonate ions having fluorine substituted at α -position as represented by the formula (f1-1) and sulfonate ions having fluorine substituted at α - and β -positions as represented by the formula (f1-2).

$$\mathbb{R}^{62}$$
— $\mathbb{C}F_2$ — \mathbb{SO}_3 -

In formula (f1-1), R^{61} is hydrogen or a C_1 - C_{20} hydrocarbyl group which may contain an ether bond, ester bond, carbonyl moiety, lactone ring, or fluorine atom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof are as will be exemplified for the hydrocarbyl group represented by R^{107} in formula (1A').

In formula (f1-2), R^{62} is hydrogen, or a C_1 - C_{30} hydrocarbyl group, C_2 - C_{30} hydrocarbylcarbonyl group, or aryloxy group, which may contain an ether bond, ester bond, carbonyl moiety or lactone ring. The hydrocarbyl group and hydrocarbyl moiety of the hydrocarbylcarbonyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof are as will be exemplified for the hydrocarbyl group represented by R^{107} in formula (1A').

Examples of the cation in the monomer from which recurring unit (f1) is derived are shown below, but not limited thereto. \mathbb{R}^4 is as defined above.

$$\mathbb{R}^{A}$$
 \mathbb{R}^{A} \mathbb{R}^{A}

$$\mathbb{R}^{A}$$
 \mathbb{R}^{A}
 \mathbb{R}^{A}
 \mathbb{R}^{A}
 \mathbb{R}^{A}

Examples of the cation in the monomer from which recurring unit (f2) or (f3) is derived are as will be exemplified for the cation in the sulfonium salt having formula (1-1).

Examples of the anion in the monomer from which recurring unit (f2) is derived are shown below, but not limited thereto. R^4 is as defined above.

$$F_{3}C \xrightarrow{F_{2}} SO_{3}^{-}$$

-continued

$$R^d$$
 O
 O
 CF_3
 CF_3

Examples of the anion in the monomer from which recurring unit (f3) is derived are shown below, but not limited thereto. \mathbb{R}^4 is as defined above.

The attachment of an acid generator to the polymer main chain is effective in restraining acid diffusion, thereby preventing a reduction of resolution due to blur by acid diffusion. Also LWR or CDU is improved since the acid generator is uniformly distributed. Where a base polymer comprising recurring units (f) is used, an acid generator of addition type (to be described later) may be omitted.

The base polymer for formulating the positive resist composition comprises recurring units (a1) or (a2) having an 60 acid labile group as essential component and additional recurring units (b), (c), (d), (e), and (f) as optional components. A fraction of units (a1), (a2), (b), (c), (d), (e), and (f) is: preferably $0 \le a1 < 1.0$, $0 \le a2 < 1.0$, 0 < a1 + a2 < 1.0, $0 \le b \le 0.9$, $0 \le c \le 0.8$, $0 \le c \le 0.8$, and $0 \le f \le 0.5$; more preferably 65 $0 \le a1 \le 0.9$, $0 \le a2 \le 0.9$, $0.1 \le a1 + a2 \le 0.9$, $0 \le b \le 0.8$, $0 \le c \le 0.8$, $0 \le c \le 0.7$, and $0 \le f \le 0.4$; and even more preferably

 $0 \le a1 \le 0.8$, $0 \le a2 \le 0.8$, $0.1 \le a1 + a2 \le 0.8$, $0 \le b \le 0.75$, $0 \le c \le 0.75$, $0 \le d \le 0.6$, 0.6, 0.6, 0.6, and $0 \le f \le 0.3$. Notably, f = f1 + f2 + f3, meaning that unit (f) is at least one of units (f1) to (3), and a1 + a2 + b + c + d + e + f = 1.0.

For the base polymer for formulating the negative resist composition, an acid labile group is not necessarily essential. The base polymer comprises recurring units (b), and optionally recurring units (c), (d), (e), and/or (f). A fraction of these units is: preferably $0 < b \le 1.0$, $0 \le c \le 0.9$, $0 \le d \le 0.8$, and $0 \le f \le 0.5$; more preferably $0.2 \le b \le 1.0$, $0 \le c \le 0.8$, $0 \le d \le 0.7$, $0 \le c \le 0.7$, and $0 \le f \le 0.4$; and even more preferably $0.3 \le b \le 1.0$, $0 \le c \le 0.75$, $0 \le d \le 0.6$, $0 \le c \le 0.6$, and $0 \le f \le 0.3$. Notably, f = f1 + f2 + f3, meaning that unit (f) is at least one of units (f1) to (3), and b + c + d + e + f = 1.0.

The base polymer may be synthesized by any desired methods, for example, by dissolving one or more monomers selected from the monomers corresponding to the foregoing recurring units in an organic solvent, adding a radical polymerization initiator thereto, and heating for polymerization. Examples of the organic solvent which can be used for polymerization include toluene, benzene, tetrahydrofuran, diethyl ether, and dioxane. Examples of the polymerization initiator used herein include 2,2'-azobisisobutyronitrile (AIBN), 2,2'-azobis(2,4-dimethylvaleronitrile), dimethyl 2,2-azobis(2-methylpropionate), benzoyl peroxide, and lauroyl peroxide. Preferably the polymerization temperature is 50 to 80° C., and the reaction time is 2 to 100 hours, more preferably 5 to 20 hours.

In the case of a monomer having a hydroxyl group, the hydroxyl group may be replaced by an acetal group susceptible to deprotection with acid, typically ethoxyethoxy, prior to polymerization, and the polymerization be followed by deprotection with weak acid and water. Alternatively, the hydroxyl group may be replaced by an acetyl, formyl, pivaloyl or similar group prior to polymerization, and the polymerization be followed by alkaline hydrolysis.

When hydroxystyrene or hydroxyvinylnaphthalene is copolymerized, an alternative method is possible. Specifically, acetoxystyrene or acetoxyvinylnaphthalene is used instead of hydroxystyrene or hydroxyvinylnaphthalene, and after polymerization, the acetoxy group is deprotected by alkaline hydrolysis, for thereby converting the polymer product to hydroxystyrene or hydroxyvinylnaphthalene. For alkaline hydrolysis, a base such as aqueous ammonia or triethylamine may be used. Preferably the reaction temperature is -20° C. to 100° C., more preferably 0° C. to 60° C., and the reaction time is 0.2 to 100 hours, more preferably 0.5 to 20 hours.

The base polymer should preferably have a weight average molecular weight (Mw) in the range of 1,000 to 500,000, and more preferably 2,000 to 30,000, as measured by GPC versus polystyrene standards using tetrahydrofuran (THF) solvent. With too low a Mw, the resist composition may become less heat resistant. A polymer with too high a Mw may lose alkaline solubility and give rise to a footing phenomenon after pattern formation.

If a base polymer has a wide molecular weight distribution or dispersity (Mw/Mn), which indicates the presence of lower and higher molecular weight polymer fractions, there is a possibility that foreign matter is left on the pattern or the pattern profile is degraded. The influences of Mw and Mw/Mn become stronger as the pattern profile becomes finer. Therefore, the base polymer should preferably have a narrow dispersity (Mw/Mn) of 1.0 to 2.0, especially 1.0 to 1.5, in order to provide a resist composition suitable for micropatterning to a small feature size.

It is understood that a blend of two or more polymers which differ in compositional ratio, Mw or Mw/Mn is acceptable.

Acid Generator

The resist composition may comprise an acid generator capable of generating a strong acid (referred to as acid generator of addition type, hereinafter). As used herein, the term"strong acid" refers to a compound having a sufficient acidity to induce deprotection reaction of an acid labile group on the base polymer in the case of a chemically amplified positive resist composition, or a compound having a sufficient acidity to induce acid-catalyzed polarity switch reaction or crosslinking reaction in the case of a chemically amplified negative resist composition. The inclusion of such an acid generator ensures that the iodized or brominated hydrocarbyl-containing carboxylic salt functions as a quencher and the inventive resist composition functions as a chemically amplified positive or negative resist composition.

The acid generator is typically a compound (PAG) capable of generating an acid upon exposure to actinic ray or radiation. Although the PAG used herein may be any compound capable of generating an acid upon exposure to high-energy radiation, those compounds capable of generating sulfonic acid, sulfonimide or sulfonmethide are preferred. Suitable PAGs include sulfonium salts, iodonium salts, sulfonyldiazomethane, N-sulfonyloxyimide, and oxime-O-sulfonate acid generators. Exemplary PAGs are described in JP-A 2008-111103, paragraphs [0122]-[0142] 30 (U.S. Pat. No. 7,537,880).

As the PAG used herein, sulfonium salts having the formula (1-1) and iodonium salts having the formula (1-2) are also preferred.

$$\begin{array}{c} R^{104} \\ \downarrow \\ I^+ \ X^- \\ \downarrow \\ R^{105} \end{array}$$

In formulae (1-1) and (1-2). R^{101} to R^{105} are each independently fluorine, chlorine, bromine, iodine, or a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom.

The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include C₁-C₂₀ alkyl groups such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl, n-hexyl, n-octyl, n-nonyl, n-decyl, undecyl, dodecyl, tri- 55 decyl, tetradecyl, pentadecyl, heptadecyl, octadecyl, nonadecyl and icosyl; C₃-C₂₀ cyclic saturated hydrocarbyl groups such as cyclopropyl, cyclopentyl, cyclohexyl, cyclopropylmethyl, 4-methylcyclohexyl, cyclohexylmethyl, norbornyl, and adamantyl: C2-C20 alkenyl groups such as vinyl, 60 propenyl, butenyl, and hexenyl; C_2 - C_{20} cyclic unsaturated aliphatic hydrocarbyl groups such as cyclohexenyl and norbornenyl; C2-C20 alkynyl groups such as ethynyl, propynyl, and butynyl; C₆-C₂₀ aryl groups such as phenyl, methylphenyl, ethylphenyl, n-propylphenyl, isopropylphe- 65 nyl, n-butylphenyl, isobutylphenyl, sec-butylphenyl, tertbutylphenyl, naphthyl, methylnaphthyl, ethylnaphthyl,

n-propylnaphthyl, isopropylnaphthyl, n-butylnaphthyl, isobuylnaphthyl, sec-butylnaphthyl, and tert-butylnaphthyl; and $\mathrm{C_7-C_{20}}$ aralkyl groups such as benzyl and phenethyl. In the foregoing groups, some hydrogen may be substituted by a moiety containing a heteroatom such as oxygen, sulfur, nitrogen or halogen, and some carbon may be replaced by a moiety containing a heteroatom such as oxygen, sulfur or nitrogen, so that the group may contain a hydroxyl, cyano, carbonyl, ether bond, ester bond, sulfonate bond, carbonate, lactone ring, sultone ring, carboxylic anhydride, or haloalkyl moiety.

 R^{101} and R^{102} may bond together to form a ring with the sulfur atom to which they are attached. Preferred rings are 15 of the structures shown below.

Herein the broken line designates an attachment to R¹⁰³. Examples of the cation in the sulfonium at having formula (1-1) are shown below, but not limited thereto.

$$\begin{array}{c} 50 \\ \hline \\ 55 \\ \hline \\ 60 \\ \hline \\ 65 \\ \hline \end{array}$$

-continued

$$S^{*}$$
 S^{*}
 S^{*}

Examples of the cation in the iodonium salt having formula (1-2) are shown below, but not limited thereto. $^{50}$

15

20

25

30

35

40

In formulae (1-1) and (1-2), X⁻ is an anion of the following formula (1A), (1B), (1C) or (1D).

$$R^{fa}$$
 — CF_2 — SO_3 -

$$R^{fb1} - CF_2 - SO_2 \tag{1B}$$

$$\begin{array}{c}
R^{\ell 2} \\
CF_{2} \\
| \\
CSO_{2} \\
| \\
SO_{2} \\
| \\
CF_{2} \\
| \\
SO_{2} \\
| \\
CF_{2} \\
| \\
R^{\ell 3}
\end{array}$$

In formula (1A), R^{fa} is fluorine or a C₁-C₄₀ hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include those exemplified later for R¹⁰⁷ in formula (1A').

Of the anions of formula (1A), an anion having the formula (1A') is preferred.

In formula (1A'), R106 is hydrogen or trifluoromethyl, 60 preferably trifluoromethyl. R^{107} is a C_1 - C_{38} hydrocarbyl group which may contain a

heteroatom. As the heteroatom, oxygen, nitrogen, sulfur and halogen atoms are preferred, with oxygen being most preferred. Of the hydrocarbyl groups represented by R¹⁰⁷, those 65 groups of 6 to 30 carbon atoms are preferred from the aspect of achieving a high resolution in forming patterns of fine feature size. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include alkyl groups such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, neopentyl, hexyl, heptyl, 2-ethylhexyl, nonyl, undecyl, tridecyl, pentadecyl, heptadecyl, and icosanyl; cyclic saturated hydrocarbyl groups such as cyclopentyl, cyclohexyl, 1-adamantyl, 2-adamantyl, 1-adamantylmethyl, norbornyl, norbornylmethyl, tricyclodecanyl, tetracyclododecanyl, tetracyclododecanylmethyl, dicyclohexyhnethyl; and unsaturated aliphatic hydrocarbyl groups such as allyl and 3-cyclohexenyl; aryl groups such as phenyl, 1-naphthyl and 2-naphthyl; and aralkyl groups such as benzyl and diphenylmethyl. In the foregoing groups, some or all hydrogen atoms may be substituted by a moiety containing a heteroatom such as oxygen, sulfur, nitrogen or halogen, and some $_{15}$ carbon may be replaced by a moiety containing a heteroatom such as oxygen, sulfur or nitrogen, so that the group may contain a hydroxyl, cyano, carbonyl, ether bond, ester bond, sulfonate bond, carbonate, lactone ring, sultone ring, carboxylic anhydride, or haloalkyl moiety. Examples of the 20 heteroatom-containing hydrocarbyl group include tetrahydrofuryl, methoxymethyl, ethoxymethyl, methylthiomethyl, acetamidemethyl, trifluoroethyl, (2-methoxyethoxy)methyl, acetoxymethyl, 2-carboxy-1-cyclohexyl, 2-oxopropyl, 4-oxo-1-adamantyl, and 3-oxocyclohexyl.

With respect to the synthesis of the sulfonium salt having an anion of formula (1A'), reference may be made to JP-A 2007-145797, JP-A 2008-106045, JP-A 2009-007327, and JP-A 2009-258695. Also useful are the sulfonium salts described in JP-A 2010-215608, JP-A 2012-041320, JP-A 2012-106986, and JP-A 2012-153644.

Examples of the anion having formula (1A) we shown below, but not limited thereto.

$$\begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \\ \\ \end{array} \end{array} \end{array} \end{array} \begin{array}{c} \begin{array}{c} \\ \\ \end{array} \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \end{array} \end{array} \begin{array}{c} \\ \end{array}$$

In formula (1B), R^{fb1} and R^{fb2} are each independently fluorine or a C_1 - C_{40} hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic, and examples thereof are as exemplified above for R^{107} . Preferably R^{fb1} and R^{fb2} are fluorine or C_1 - C_4 straight fluorinated alkyl groups. Also, R^{fb1} and R^{fb2} may bond together to form a ring with the linkage: $-CF_2$ - $-SO_2N^-$ - $-SO_2$ - $-CF_2$ - to which they are attached. It is preferred that a combination of R^{fb1} and R^{fb2} be a fluorinated ethylene or fluorinated propylene group.

In formula (1C), R^{fc1} , R^{fc2} and R^{fc3} are each independently fluorine or a C_1 - C_{40} hydrocarbyl group which may

25

contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic, and examples thereof are as exemplified above for R^{107} . Preferably R^{fc1} , R^{fc2} and R^{fc3} are fluorine or $C_1\text{-}C_4$ straight fluorinated alkyl groups. Also, R^{fc1} and R^{fc2} may bond together to form a ring with the linkage: —CF $_2$ —SO $_2$ — C 2 —To which they are attached. It is preferred that a combination of R^{fc1} and R^{fc2} be a fluorinated ethylene or fluorinated propylene group.

In formula (1D), R^{fd} is a C_1 - C_{40} hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic, and examples thereof are as exemplified above for R^{107} .

With respect to the synthesis of the sulfonium salt having ¹⁵ an anion of formula (1D), reference may be made to JP-A 2010-215608 and JP-A 2014-133723.

Examples of the anion having formula (1D) are shown below, but not limited thereto.

$$\begin{array}{c} O \\ CF_3 \\ SO_3^- \\ CF$$

Notably, the compound having the anion of formula (1D) does not have fluorine at the α -position relative to the sulfo group, but two trifluoromethyl groups at the β -position. For this reason, it has a sufficient acidity to sever the acid labile groups in the base polymer. Thus the compound is an effective PAG.

Another preferred PAG is a compound having the formula $_{35}$ (2).

$$R^{201} - S^{+} - R^{203} - L^{A} - \begin{pmatrix} X^{C} & X^{A} \\ C & C \\ X^{D} & X^{B} \end{pmatrix} = SO_{3}^{-}$$
(2)

In formula (2), R²⁰¹ and R²⁰² are each independently a C₁-C₃₀ hydrocarbyl group which may contain a heteroatom. R²⁰³ is a C₁-C₃₀ hydrocarbylene group which may contain a heteroatom. Any two of R²⁰¹, R²⁰² and R²⁰³ may bond together to form a ring with the sulfur atom to which they are attached. Examples of the ring are as exemplified above for the ring that R¹⁰¹ and R¹⁰² in formula (1-1), taken together, form with the sulfur atom to which they are attached.

form with the sulfur atom to which they are attached.

The hydrocarbyl groups R²⁰¹ and R²⁰² may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include alkyl groups such as methyl, ethyl, propyl, isopropyl, n-butyl, sec-butyl, tert-butyl, n-pentyl, tert-pentyl, n-hexyl, n-octyl, 2-ethylhexyl, n-nonyl, and n-decyl; cyclic saturated hydrocarbyl groups such as cyclopentyl, cyclohexyl, cyclopentylmethyl, cyclopentylethyl, cyclopentylbutyl, cyclohexylmethyl, cyclohexylethyl, cyclohexylbutyl, norbornyl, tricyclo[5.2.1.0^{2,6}]decanyl, and adamantyl: aryl groups such as phenyl, naphthyl, and anthracenyl. In the foregoing groups, some hydrogen may be substituted by a moiety containing a heteroatom such as oxygen, sulfur, nitrogen or halogen, and some carbon may be replaced by a moiety containing a heteroatom such as oxygen, sulfur or nitrogen, so that the group may contain a hydroxyl, cyano,

carbonyl, ether bond, ester bond, sulfonate bond, carbonate, lactone ring, sultone ring, carboxylic anhydride or haloalkyl moiety.

The hydrocarbylene group R²⁰³ may be saturated or unsaturated and straight, branched or cyclic. Examples 5 thereof include alkanediyl groups such as methylene, ethvlene, propane-1,3-divl, butane-1,4-divl, pentane-1,5-divl, hexane-1,6-diyl, heptane-1,7-diyl, octane-1,8-diyl, nonane-1,9-diyl, decane-1,10-diyl, undecane-1,11-diyl, dodecane-1, 12-diyl, tridecane-1,13-diyl, tetradecane-1,14-diyl, pentadecane-1,15-diyl, hexadecane-1,16-diyl, and heptadecane-1, 17-diyl; cyclic saturated hydrocarbylene groups such as cyclopentanediyl, cyclohexanediyl, norbornanediyl and adamantanediyl; and arylene groups such as phenylene, meth- $_{15}$ ylphenylene, ethylphenylene, n-propylphenylene, isopropylphenylene, n-butylphenylene, isobutylphenylene, secbutyphenylene, tert-butylphenylene, naphthylene, methylnaphthylene, ethylnaphthylene, n-propylnaphthyln-butynaphthylene, 20 isopropylnaphthylene, ene. isobutylnaphthylene, sec-butylnaphthylene, and tertbutylnaphthylene. In these groups, some or all of the hydrogen atoms may be substituted by a moiety containing a heteroatom such as oxygen, sulfur, nitrogen or halogen, or some carbon may be replaced by a moiety containing a 25 heteroatom such as oxygen, sulfur or nitrogen, so that the group may contain a hydroxyl, cyano, carbonyl, ether bond, ester bond, sulfonate bond, carbonate, lactone ring, sultone ring carboxylic anhydride or haloalkyl moiety. Of the heteroatoms, oxygen is preferred.

In formula (2), L^4 is a single bond, ether bond or a C_1 - C_{20} hydrocarbylene group which may contain a heteroatom. The hydrocarbylene group may be saturated or unsaturated and straight, bunched or cyclic. Examples thereof are as exemplified above for R^{203} .

In formula (2), X^A , X^B , X^C and X^D are each independently hydrogen, fluorine or trifluoromethyl, with the proviso that at least one of X^A , X^B , X^C and X^D is fluorine or trifluoromethyl, and k is an integer of 0 to 3.

Of the PAGs having formula (2), those having formula $_{40}$ (2') are preferred.

$$(R^{301})_{x} \qquad (R^{303})_{z} \qquad H \qquad F \qquad (2')$$

$$L^{A} - C - C - SO_{3} \qquad R^{HF} \qquad F$$

$$(R^{302})_{y} \qquad (R^{302})_{y} \qquad (R^{302$$

In formula (2'), L^4 is as defined above. R^{HF} is hydrogen or trifluoromethyl, preferably trifluoromethyl. R^{301} , R^{302} 55 and R^{303} are each independently hydrogen or a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof are as exemplified above for R^{107} in formula (1A'). The subscripts x and 60 y are each independently an integer of 0 to 5, and z is an integer of 0 to 4.

Of the foregoing PAGs, those having an anion of formula (1A') or (1D) are especially preferred because of reduced acid diffusion and high solubility in the resist solvent. Also 65 those having an anion of formula (2') are especially preferred because of extremely reduced acid diffusion.

118

Also a sulfonium or iodonium salt having an iodized or brominated aromatic ring-containing anion may be used as the PAG. Suitable are sulfonium and iodonium salts having the formulae (3-1) and (3-2).

$$\begin{bmatrix} (X^{BI})_s & Rf^1 & R^{402} \\ \vdots & \vdots & \vdots \\ (R^{401})_t & Rf^3 & Rf^4 \end{bmatrix}$$
(3-1)

In formulae (3-1) and (3-2), r is an integer of to 3, s is an integer of 1 to 5, and t is an integer of 0 to 3, and $1 \le s+t \le 5$. Preferably, s is an integer of 1 to 3, more preferably 2 or 3, and t is an integer of 0 to 2.

 X^{BI} is iodine or bromine, and may be the same or different when r and/or s is 2 or more.

L¹ is a single bond, ether bond, ester bond, or a C₁-C₆ saturated hydrocarbylene group which may contain an ether bond or ester bond. Ie saturated hydrocarbylene group may be straight, branched or cyclic.

 $\rm L^2$ is a single bond or a $\rm C_1\text{-}C_{20}$ divalent linking group when r=1, or a $\rm C_1\text{-}C_{20}$ (r+1)-valent linking group when r=2 or 3, the linking group optionally containing an oxygen, sulfur or nitrogen atom.

R⁴⁰¹ is a hydroxyl group, carboxyl group, fluorine, chlorine, bromine, amino group, or a C1-C20 saturated hydrocarbyl, C₁-C₂₀ saturated hydrocarbyloxy, C₂-C₁₀ saturated hydrocarbyloxycarbonyl, C2-C20 saturated hydrocarbylcarbonyloxy or C₁-C₂₀ saturated hydrocarbylsulfonyloxy group, which may contain fluorine, chlorine, bromine, hydroxyl, amino or ether bond, or $-NR^{401A}$ —C(=O)—45 R^{401B} or $-NR^{401A}$ —C(=O)—O— R^{401B} . R^{401A} is hydrogen or a C₁-C₆ saturated hydrocarbyl group which may contain halogen, hydroxyl, C_1 - C_6 saturated hydrocarbyloxy. $\rm C_2$ - $\rm C_6$ saturated hydrocarbylcarbonyl or $\rm C_2$ - $\rm C_6$ saturated hydrocarbylcarbonyloxy moiety. $\rm R^{401B}$ is a $\rm C_1$ - $\rm C_{16}$ aliphatic $_{50}\,$ hydrocarbyl or $\mathrm{C_{6}\text{-}C_{12}}$ aryl group, which may contain halogen, hydroxyl, C_1 - C_6 saturated hydrocarbyloxy, C_2 - C_6 saturated hydrocarbykarbonyl or C2-C6 saturated hydrocarbylcarbonyloxy moiety. The aliphatic hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. The saturated hydrocarbyl, saturated hydrocarbyloxy, saturated hydrocarbyloxycarbonyl, saturated hydrocarbylcarbonyl, and saturated hydrocarbylcarbonyloxy groups may be straight, branched or cyclic. Groups R⁴⁰¹ may be the same or different when r and/or s is 2 or more. Of these, R^{401} is preferably hydroxyl, $-NR^{401A}-C(-0)-R^{401B}$, preferably —NR⁴⁰¹A--C(=O)—O— R^{401B} , fluorine, chlorine, bromine, methyl or methoxy.

In formulae (3-1) and (3-2), Rf¹ to Rf⁴ are each independently hydrogen, fluorine or trifluoromethyl, at least one of Rf¹ to RF⁴ is fluorine or trifluoromethyl, or Rf¹ and Rf², taken together, may form a carbonyl group. Preferably, both Rf⁵ and Rf⁴ are fluorine.

 $R^{402},\,R^{403},\,R^{404},\,R^{405}$ and R^{406} are each independently a $C_1\text{-}C_{20}$ hydrocarbyl group which may contain a heteroatom. The hydrocarbyl group may be saturated or unsaturated and straight, branched or cyclic. Examples thereof include $C_1\text{-}C_{20}$ alkyl, $C_3\text{-}C_{20}$ cycloalkyl, $C_2\text{-}C_{20}$ alkenyl, $C_2\text{-}C_{20}$ alkynyl, $C_6\text{-}C_{20}$ aryl, and $C_7\text{-}C_{20}$ aralkyl groups. In these groups, some or all of the hydrogen atoms may be substituted by hydroxyl, carboxyl, halogen, cyano, nitro, mercapto, sultone, sulfone, or sulfonium salt-containing moieties, and some carbon may be replaced by an ether bond, ester bond, carbonyl moiety, amide bond, carbonate moiety or sulfonic acid ester bond. Any two of $R^{402},\,R^{403}$ and R^{404} may bond together to form a ring with the sulfur atom to which they are attached. Exemplary rings are the same as described above for the ring that R^{101} or and R^{102} in formula 15 (1-1), taken together, form with the sulfur atom to which they are attached.

Examples of the cation in the sulfonium salt having formula (3-1) include those exemplified above as the cation in the sulfonium salt having formula (1-1). Examples of the ²⁰ cation in the iodonium salt having formula (3-2) include those exemplified above as the cation in the iodonium salt having formula (1-2).

Examples of the anion in the onium salts having formulae (3-1) and (3-2) are shown below, but not limited thereto. ²⁵ Herein X^{BI} is as defined above.

$$X^{BI}$$
 CF_3
 $F_2C - SO_3$
 $T_2C - SO_3$

-continued

$$X^{BI}$$
 Y^{BI}
 $Y^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 O
 CF_3
 S_2
 S_3
 S_4
 S_4
 S_5
 S_5
 S_6
 S_7
 S_7
 S_7
 S_8
 S_7
 S_8
 $S_$

$$X^{BI}$$
 CF_3
 F_2C
 SO_3
 SO

$$X^{BI}$$
 CF_3
 CF_3
 $F_2C - SO_3$
 $F_2C - SO_3$
 $F_2C - SO_3$

-continued

$$X^{BI}$$
 O
 CF_3
 F_2C
 SO_3
 SO_3
 SO_4
 SO_4
 SO_5
 S

$$X^{BI}$$
 CF_3
 F_2C
 SO_3

-continued

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}

$$X^{BI}$$
 X^{BI}
 X

-continued

$$X^{BI}$$
-continued

 X^{BI}
-continued

 X^{BI}
 X^{BI}

-continued

-continued

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 X^{BI}
 X

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 X^{BI}
 X

$$\begin{array}{c|c}
\mathbf{US} & \mathbf{1} \\
\mathbf{131} \\
-\text{continued} \\
X^{BI} \\
X^{BI} \\
Y^{BI} \\$$

$$F_{2}C$$
 SO_{3}
 X^{BI}
 X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 X^{BI}
 X

$$X^{BI}$$
 X^{BI}
 X

$$X^{BI}$$

$$X$$

Tooltinued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 Y^{BI}
 $Y^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 F_2C
 SO_3

139
-continued

$$CBI$$
 CBI
 CBI

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$
 F_2C
 SO_3

T41

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued
$$X^{BI}$$
 X^{BI} X^{BI}

-continued CF_3

 X^{BI} O CF_3 SO_3 15

 $X^{BI} \longrightarrow CF_3$ $X^{BI} \longrightarrow CF_3$ $X^{BI} \longrightarrow CF_3$ SO_3^- 30

 $O \longrightarrow O \longrightarrow CF_3$ $O \longrightarrow CF_3$ $SO_3^ X^{BI}$

 $O \longrightarrow O \longrightarrow CF_3 \longrightarrow SO_3$ $O \longrightarrow CF_3 \longrightarrow SO_3$ $SO_3 \longrightarrow SO_3$

55

 $\begin{array}{c} X^{BI} \\ O \\ O \\ \end{array} \begin{array}{c} CF_3 \\ SO_3 \end{array}$

-continued

O

O

CF₃
SO₃
SO₃ CF_3 CF_3

 X^{BI} O CF_3 SO_3

 X^{BI} O O O CF_3 SO_3

-continued

$$X^{BI}$$
 CF_3
 CF_3
 SO_3

15

 CF_3
 SO_3

15

 CF_3
 SO_3
 SO_3

$$X^{BI}$$
 X^{BI}
 X^{B

-continued

$$X^{BI}$$

$$X^$$

-continued

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}

-continued
$$X^{BI}$$
 X^{BI} X^{BI}

$$X^{BI}$$
 X^{BI}
 X

$$X^{BI}$$
 X^{BI}
 X

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}

$$X^{BI}$$
 X^{BI}
 X^{BI}

$$X^{BI}$$
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}
 X^{BI}

-continued
$$X^{BI}$$
 X^{BI} X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

-continued

5

HN

$$F_2C$$
— SO_3 -

15

HN

20

$$X^{BI}$$
 X^{BI}
 X

$$X^{BI}$$

$$X$$

$$F_{2}C \longrightarrow SO_{3}^{-}$$

$$X^{BI} \longrightarrow X^{BI}$$

$$60$$

-continued

-continued

$$X^{BI}$$
 X^{BI}
 X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

HO

$$X^{BI}$$
 X^{BI}
 X^{BI}

-continued
$$X^{BI}$$
 X^{BI} X^{BI}

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

$$X^{BI}$$

$$X$$

-continued

$$X^{BI}$$
 X^{BI}
 $X^$

-continued
$$X^{BI}$$
 X^{BI} X^{BI}

$$X^{BI}$$
 X^{BI}
 X^{B

When used, the acid generator of addition type is preferably added in an amount of 0.1 to 50 parts, and more preferably 1 to 40 parts by weight per 100 parts by weight of the base polymer. The resist composition functions as a chemically amplified resist composition when the base poly-15 mer includes recurring units (f) and/or the resist composition contains the acid generator of addition type.

Organic Solvent

An organic solvent may be added to the resist composition. The organic solvent used herein is not particularly 20 limited as long as the foregoing and other components are soluble therein. Examples of the organic solvent are described in JP-A 2008-111103, paragraphs [0144]-[0145] (U.S. Pat. No. 7,537,880). Exemplary solvents include ketones such as cyclohexanone, cyclopentanone, methyl-2-25 n-pentyl ketone and 2-heptanone; alcohols such as 3-methoxybutanol, 3-methyl-3-methoxybutanol, 1-methoxy-2-propanol, 1-ethoxy-2-propanol and diacetone alcohol (DAA); ethers such as propylene glycol monomethyl ether (PGME), ethylene glycol monomethyl ether, 30 propylene glycol monoethyl ether, ethylene glycol monoethyl ether, propylene glycol dimethyl ether, and diethylene glycol dimethyl ether; esters such as propylene glycol monomethyl ether acetate (PGMEA), propylene glycol monoethyl ether acetate, ethyl lactate, ethyl pyruvate, butyl 35 acetate, methyl 3-methoxypropionate, ethyl 3-ethoxypropionate, tert-butyl acetate, tert-butyl propionate, and propylene glycol mono-tert-butyl ether acetate; and lactones such as γ-butyrolactone, which may be used alone or in admix-

40 The organic solvent is preferably added in an amount of 100 to 10,000 parts, and more preferably 200 to 8,000 parts by weight per 100 parts by weight of the base polymer.

Other Components

With the foregoing components, other components such 45 as a surfactant, dissolution inhibitor, and crosslinker may be blended in any desired combination to formulate a chemically amplified positive or negative resist composition. This positive or negative resist composition has a very high sensitivity in that the dissolution rate in developer of the 50 base polymer in exposed areas is accelerated by catalytic reaction. In addition, the resist film has a high dissolution contrast, resolution, exposure latitude, and process adaptability, and provides a good pattern profile after exposure, and minimal proximity bias because of restrained acid 55 diffusion. By virtue of these advantages, the composition is fully useful in commercial application and suited as a pattern-forming material for the fabrication of VLSIs.

Exemplary surfactants are described in JP-A 2008-111103, paragraphs [0165]-[0166]. Inclusion of a surfactant 60 may improve or control the coating characteristics of the resist composition. While the surfactant may be used alone or in admixture, it is preferably added in an amount of 0.0001 to 10 parts by weight per 100 parts by weight of the base polymer.

In the case of positive resist compositions, inclusion of a dissolution inhibitor may lead to an increased difference in dissolution rate between exposed and unexposed areas and

a further improvement in resolution. The dissolution inhibitor which can be used herein is a compound having at least two phenolic hydroxyl groups on the molecule, in which an average of from 0 to 100 mol % of all the hydrogen atoms on the phenolic hydroxyl groups are replaced by acid labile 5 groups or a compound having at least one carboxyl group on the molecule, in which an average of 50 to 100 mol % of all the hydrogen atoms on the carboxyl groups are replaced by acid labile groups, both the compounds having a molecular weight of 100 to 1,000, and preferably 150 to 800. Typical 10 are bisphenol A, trisphenol, phenolphthalein, cresol novolac, naphthalenecarboxylic acid, adamantanecarboxylic acid, and cholic acid derivatives in which the hydrogen atom on the hydroxyl or carboxyl group is replaced by an acid labile group, as described in U.S. Pat. No. 7,771,914 (JP-A 2008- 15 122932, paragraphs [0155]-[0178]).

In the positive resist composition, the dissolution inhibitor is preferably added in an amount of 0 to 50 parts, more preferably 5 to 40 parts by weight per 100 parts by weight of the base polymer. The dissolution inhibitor may be used 20 alone or in admixture.

In the case of negative resist compositions, a negative pattern may be formed by adding a crosslinker to reduce the dissolution rate of a resist film in exposed area. Suitable crosslinkers which can be used herein include epoxy compounds, melamine compounds, glunamine compounds, glycoluril compounds and urea compounds having substituted thereon at least one group selected from among methylol, alkoxymethyl and acyloxymethyl groups, isocyanate compounds, azide compounds, and compounds having a double 30 bond such as an alkenyl ether group. These compounds may be used as an additive or introduced into a polymer side chain as a pendant. Hydroxy-containing compounds may also be used as the crosslinker. The crosslinker may be used alone or in admixture.

Of the foregoing crosslinkers, examples of the epoxy compound include tris(2,3-epoxypropyl) isocyanurate, trimethylohmethane triglycidyl ether, trimethylolpropane triglycidyl ether, and triethylolethane triglycidyl ether. Examples of the melamine compound include hexamethylol melamine, 40 hexamethoxymethyl melamine, hexamethylol melamine compounds having 1 to 6 methylol groups methoxymethylated and mixtures thereof hexamethoxyethyl melamine, hexaacyloxymethyl melamine, hexamethylol melamine compounds having 1 to 6 methylol groups acyloxymethyl- 45 ated and mixtures thereof. Examples of the guanamine compound include tetramethylol guanamine, tetramethoxymethyl guanamine, tetramethylol guanamine compounds having 1 to 4 methylol groups methoxymethylated and mixtures thereof, tetramethoxyethyl guanamine tetraa- 50 cyloxyguanamine tetramethylol guanamine compounds having 1 to 4 methylol groups acyloxymethylated and mixtures thereof. Examples of the glycoluril compound include tetramethylol glycoluril, tetramethoxyglycoluril, tetramethoxymethyl glycoluril, tetramethylol glycoluril 55 compounds having 1 to 4 methylol groups methoxymethylated and mixtures thereof tetramethylol glycoluril compounds having 1 to 4 methylol groups acyloxymethylated and mixtures thereof. Examples of the urea compound include tetramethylolurea, tetramethoxymethyl urea, tetram- 60 ethylol urea compounds having 1 to 4 methylol groups methoxymethylated and mixtures thereof, and tetramethoxyethyl urea.

Suitable isocyanate compounds include tolylene diisocyanate, diphenylmethane diisocyanate, hexamethylene diisocyanate and cyclohexane diisocyanate. Suitable azide compounds include 1,1'-biphenyl-4,4'-bisazide, 4,4'-

methylidenebisazide, and 4,4'-oxybisazide. Examples of the alkenyl ether group-containing compound include ethylene glycol divinyl ether, triethylene glycol divinyl ether, 1,2-propanediol divinyl ether, 1,4-butanediol divinyl ether, tetramethylene glycol divinyl ether, neopentyl glycol divinyl ether, trimethylol propane trivinyl ether, hexanediol divinyl ether, 1,4-cyclohexanediol divinyl ether, pentaerythritol trivinyl ether, pentaerythritol tetravinyl ether, sorbitol tetravinyl ether, sorbitol pentavinyl ether, and trimethylol propane trivinyl ether.

In the negative resist composition, the crosslinker is preferably added in an amount of 0 to 50 parts, more preferably 1 to 40 parts by weight per 100 parts by weight of the base polymer.

In the resist composition of the invention, a quencher other than the inventive iodized or brominated hydrocarbylcontaining carboxylic salt may be blended. The other quencher is typically selected from conventional basic compounds. Conventional basic compounds include primary, secondary, and tertiary aliphatic amines, mixed amines, aromatic amines, heterocyclic amines, nitrogen-containing compounds with carboxyl group, nitrogen-containing compounds with sulfonyl group, nitrogen-containing compounds with hydroxyl group, nitrogen-containing compounds with hydroxyphenyl group, alcoholic nitrogencontaining compounds, amide derivatives, derivatives, and carbamate derivatives. Also included are primary, secondary, and tertiary amine compounds, specifically amine compounds having a hydroxyl group, ether bond, ester bond, lactone ring, cyano group, or sulfonic acid ester bond as described in JP-A 2008-111103, paragraphs [0146]-[0164], and compounds having a carbamate group as described in JP 3790649. Addition of a basic compound may be effective for further suppressing the diffusion rate of acid 35 in the resist film or correcting the pattern profile.

Onium salts such as sulfonium salts, iodonium salts and ammonium salts of sulfonic acids which are not fluorinated at $\alpha\text{-position}$ as described in U.S. Pat. No. 8,795,942 (JP-A 2008-158339) and similar onium salts of carboxylic acid may also be used as the other quencher. While an $\alpha\text{-fluorinated}$ sulfonic acid, sulfonimide, and sulfonemethide are necessary to deprotect the acid labile group of carboxylic acid ester, an $\alpha\text{-non-fluorinated}$ sulfonic acid and a carboxylic acid are released by salt exchange with an $\alpha\text{-non-fluorinated}$ onium salt. An $\alpha\text{-non-fluorinated}$ sulfonic acid and a carboxylic acid function as a quencher because they do not induce deprotection reaction.

Also useful are quenchers of polymer type as described in U.S. Pat. No. 7,598,016 (JP-A 2008-239918). The polymeric quencher segregates at the resist surface after coating and thus enhances the rectangularity of resist pattern. When a protective film is applied as is often the case in the immersion lithography, the polymeric quencher is also effective for preventing a film thickness loss of resist pattern or rounding of pattern top.

The other quencher is preferably added in an amount of 0 to 5 parts, more preferably 0 to 4 parts by weight per 100 parts by weight of the base polymer. The other quencher may be used alone or in admixture.

To the resist composition, a water repellency improver may also be added for improving the water repellency on surface of a resist film as spin coated. The water repellency improver may be used in the topcoatless immersion lithography. Suitable water repellency improvers include polymers having a fluoroalkyl group and polymers having a specific structure with a 1,1,1,3,3,3-hexafluoro-2-propanol residue and me described in JP-A 2007-297590 and JP-A

2008-111103, for example. The water repellency improver to be added to the resist composition should be soluble in the alkaline developer and organic solvent developer. The water repellency improver of specific structure with a 1,1,1,3,3,3-hexafluoro-2-propanol residue is well soluble in the developer. A polymer having an amino group or amine salt copolymerized as recurring units may serve as the water repellent additive and is effective for preventing evaporation of acid during PEB, thus preventing any hole pattern opening failure after development. The water repellency improver may be used alone or in admixture. An appropriate amount of the water repellency improver is 0 to 20 parts, more preferably 0.5 to 10 parts by weight per 100 parts by weight of the base polymer.

Also, an acetylene alcohol may be blended in the resist 15 composition. Suitable acetylene alcohols are described in JP-A 2008-122932, paragraphs [0179]-[0182]. An appropriate amount of the acetylene alcohol blended is 0 to 5 parts by weight per 100 parts by weight of the base polymer. Pattern Forming Process 20

The resist composition is used in the fabrication of various integrated circuits.

Pattern formation using the resist composition may be performed by well-known lithography processes. The process generally involves coating, exposure, and development. 25 If necessary, any additional steps may be added.

For example, the resist composition is first applied onto a substrate on which an integrated circuit is to be formed (e.g., Si, SiO₂, SiN, SiON, TiN, WSi, BPSG, SOG, or organic antireflective coating) or a substrate on which a mask circuit 30 is to be formed (e.g., Cr, CrO, CrON, MoSi₂, or SiO₂) by a suitable coating technique such as spin coating, roll coating, flow coating, dipping, spraying or doctor coating. The coating is prebaked on a hot plate at a temperature of 60 to 150° C. for 10 seconds to 30 minutes, preferably at 80 to 35 120° C. for 30 seconds to 20 minutes. The resulting resist film is generally 0.1 to 2 μ m thick.

The resist film is then exposed to a desired pattern of high-energy radiation such as UV, deep-UV, EB, EUV, x-ray, soft x-ray, excimer laser light, γ-ray or sychrotron radiation. 40 When UV, deep-UV, EUV, x-ray, soft x-ray, excimer laser light, γ-ray or synchrotron radiation is used as the highenergy radiation, the resist film is exposed thereto through a mask having a desired pattern in a dose of preferably about 1 to 200 mJ/cm², more preferably about 10 to 100 mJ/cm². 45 When EB is used as the high-energy radiation, the resist film is exposed thereto through a mask having a desired pattern or directly in a dose of preferably about 0.1 to 100 μC/cm², more preferably about 0.5 to 50 μC/cm². It is appreciated that the inventive resist composition is suited in micropat- 50 terning using KrF excimer laser, ArF excimer laser, EB, EUV, x-ray, soft x-ray, γ-ray or synchrotron radiation, especially EB or EUV.

After the exposure, the resist film may be baked (PEB) on a hotplate or in an oven at 30 to 150° C. for 10 seconds to 55 30 minutes, preferably at 50 to 120° C. for 30 seconds to 20 minutes.

After the exposure or PEB, the resist film is developed in a developer in the form of an aqueous base solution for 3 seconds to 3 minutes, preferably 5 seconds to 2 minutes by 60 conventional techniques such as dip, puddle and spray techniques. A typical developer is a 0.1 to 10 wt %, preferably 2 to 5 wt % aqueous solution of tetramethylammonium hydroxide (TMAH), tetraethylammonium hydroxide (TEAH), tetrapropylammonium hydroxide 65 (TPAH), or tetrabutylammonium hydroxide (TBAH). In the case of positive resist, the resist film in the exposed area is

dissolved in the developer whereas the resist film in the unexposed area is not dissolved. In this way, the desired positive pattern is formed on the substrate. Inversely in the case of negative resist, the exposed area of resist film is insolubilized and the unexposed area is dissolved in the developer.

In an alternative embodiment, a positive resist composition comprising a base polymer having an acid labile group is used to form a negative pattern via organic solvent development. The developer used herein is preferably selected from among 2-octanone, 2-nonanone, 2-heptanone, 3-heptanone, 4-heptanone, 2-hexanone, 3-hexanone, diisobutyl ketone, methylcyclohexanone, acetophenone, methylacetophenone, propyl acetate, butyl acetate, isobutyl acetate, pentyl acetate, butenyl acetate, isopentyl acetate, propyl formate, butyl formate, isobutyl formate, pentyl formate, isopentyl formate, methyl valerate, methyl pentenoate, methyl crotonate, ethyl crotonate, methyl propionate, ethyl propionate, ethyl 3-ethoxypropionate, methyl 20 lactate, ethyl lactate, propyl lactate, butyl lactate, isobutyl lactate, pentyl lactate, isopentyl lactate, methyl 2-hydroxyisobutyrate, ethyl 2-hydroxyisobutyrate, methyl benzoate, ethyl benzoate, phenyl acetate, benzyl acetate, methyl phenylacetate, benzyl formate, phenylethyl formate, methyl 3-phenylpropionate, benzyl propionate, ethyl phenylacetate, and 2-phenylethyl acetate, and mixtures thereof.

At the end of development, the resist film is rinsed. As the rinsing liquid, a solvent which is miscible with the developer and does not dissolve the resist film is preferred. Suitable solvents include alcohols of 3 to 10 carbon atoms, ether compounds of 8 to 12 carbon atoms, alkanes, alkenes, and alkynes of 6 to 12 carbon atoms, and aromatic solvents. Specifically, suitable alcohols of 3 to 10 carbon atoms include n-propyl alcohol, isopropyl alcohol, 1-butyl alcohol, 2-butyl alcohol, isobutyl alcohol, t-butyl alcohol, 1-pentanol, 2-pentanol, 3-pentanol, t-pentyl alcohol, neopentyl alcohol, 2-methyl-1-butanol, 3-methyl-1-butanol, 3-methyl-3-pentanol, cyclopentanol, 1-hexanol, 2-hexanol, 3-hexanol, 2,3-dimethyl-2-butanol, 3,3-dimethyl-1-butanol, 3,3-dimethyl-2-butanol, 2-ethyl-1-butanol, 2-methyl-1-pentanol, 2-methyl-2-pentanol, 2-methyl-3-pentanol, 3-methyl-1-pentanol, 3-methyl-2-pentanol, 3-methyl-3-pentanol, 4-methyl-1-pentanol, 4-methyl-2-pentanol, 4-methyl-3-pentanol, cyclohexanol and 1-octanol. Suitable ether compounds of 8 to 12 carbon atoms include di-n-butyl ether, diisobutyl ether, di-s-butyl ether, di-n-pentyl ether, diisopentyl ether, di-spentyl ether, di-t-pentyl ether, and di-n-hexyl ether. Suitable alkanes of 6 to 12 carbon atoms include hexane, heptane, octane, nonane, decane, undecane, dodecane, methylcyclopentane, dimethylcyclopentane, cyclohexane, methylcyclohexane, dimethylcyclohexane, cycloheptane, cyclooctane, and cyclononane. Suitable alkenes of 6 to 12 carbon atoms include hexene, heptene, octene, cyclohexene, methylcyclodimethylcyclohexene, hexene. cycloheptene, cyclooctene. Suitable alkynes of 6 to 12 carbon atoms include hexyne, heptyne, and octyne. Suitable aromatic solvents include toluene, xylene, ethylbenzene, isopropylbenzene, t-butylbenzene and mesitylene. The solvents may be used alone or in admixture.

Rinsing is effective for minimizing the risks of resist pattern collapse and defect formation. However, rinsing is not essential. If rinsing is omitted, the amount of solvent used may be reduced.

A hole or trench pattern after development may be shrunk by the thermal flow, RELACS® or DSA process. A hole pattern is shrunk by coating a shrink agent thereto, and baking such that the shrink agent may undergo crosslinking

at the resist surface as a result of the acid catalyst diffusing from the resist layer during bake, and the shrink agent may attach to the sidewall of the hole pattern. The bake is preferably at a temperature of 70 to 180° C., more preferably 80 to 170° C. for a time of 10 to 300 seconds. The extra shrink agent is stripped and the hole pattern is shrunk.

EXAMPLES

Examples of the invention are given below by way of illustration and not by way of limitation. The abbreviation "pbw" is parts by weight.

Quenchers 1 to 24 used in resist compositions have the structure, shown below. Quenchers 1 to 24 were prepared by mixing equi-molar amounts of an iodized or brominated hydrocarbyl-containing carboxylic acid priding the anion shown below and a 2,5,8,9-tetraaza-1-phosphabicyclo [3.3.3]undecane, biguanide or phosphazene compound providing the cation shown below in methanol, and evaporating off the methanol.

Quencher 1
$$\stackrel{25}{\sim}$$
 $\stackrel{\text{Quencher 1}}{\sim}$ $\stackrel{25}{\sim}$ $\stackrel{\text{Quencher 1}}{\sim}$ $\stackrel{25}{\sim}$ $\stackrel{\text{Quencher 1}}{\sim}$ $\stackrel{25}{\sim}$

Quencher 2
$$_{35}$$

NH HN

N+H

40

Quencher 6
$$I \longrightarrow O^{-} \longrightarrow N^{NH_2} \longrightarrow N^{+}H_2$$

Quencher 11

-continued

-continued

Quencher 13
$$\begin{array}{c}
 & 30 \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 & & \\
 &$$

Quencher 14
$$_{45}$$
 $_{F}$

Quencher 14 $_{45}$
 $_{N}$
 $_{N}$

Quencher 15
$$\begin{array}{c} \text{Quencher 15} \\ \text{NH}_2 & \text{NH}_2 \\ \text{O} & \text{HN} & \text{N}^{\dagger}\text{H}_2 \end{array}$$

Quencher 17
$$NH_2$$
 NH_2 N^+H_2 N^+H_2

$$\begin{array}{c} & & \text{Quencher 18} \\ & & \text{NH}_2 & \text{NH}_2 \\ & & \text{N}^{\dagger}\text{H}_2 \\ & & \text{Br} & & \\ & & \text{Br} & & \\ & & & \text{Pr} & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ &$$

Quencher 19
$$\begin{array}{c} & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ &$$

Quencher 20
$$I \longrightarrow I \longrightarrow I$$

$$I \longrightarrow$$

Quencher 21

$$V_{N}$$
 V_{N}
 V_{N}

-continued

Quencher 22
$$\begin{array}{c} I \\ O \\ F_2C \\ O \end{array}$$

Quencher 23

Quencher 24

35

SYNTHESIS EXAMPLE

Synthesis of Base Polymers (Polymers 1 to 4)

Base polymers were prepared by combining suitable monomers, effecting copolymerization reaction thereof in tetrahydrofuran (THF) solvent, pouring the reaction Solution into Methanol for crystallization, repeatedly washing with hexane, isolation, and drying. The resulting polymers, designated Polymers 1 to 4, were analyzed for composition by ¹H-NMR Spectroscopy, and for Mw and Mw/Mn by GPC versus polystyrene standards using THF solvent.

Mw = 8,600Mw/Mn = 1.73

-continued
$$F_2$$
 SO_3 CF_3 $M_W = 8,900$

Mw/Mn = 1.89 Polymer 3

$$M_{W} = 7,600$$
 $M_{W}/M_{B} = 1.73$

20

35

PAG 2

-continued

-continued

$$Mw = 6,900$$

 $Mw/Mn = 1.62$

Examples 1 to 27 and Comparative Examples 1 to 7

Preparation of Resist Compositions

Resist compositions in solution form were prepared by dissolving components in a solvent in accordance with the recipe shown in Tables 1 to 3, and filtering through a filter having a pore size of 0.2 μm . The solvent contained 100 ppm of surfactant FC-4430 (3M). The resist compositions of Examples 1 to 26 and Comparative Examples 1 to 6 were of positive tone, while the resist compositions of Example 27 $_{30}$ and Comparative Example 7 were of negative tone.

The components in Tables 1 to 3 are as identified below. Polymers 1 to 4 of the above structural formulae Organic Solvents:

PGMEA (propylene glycol monomethyl ether acetate) CyH (cyclohexanone)

PGME (propylene glycol monomethyl ether)
DAA (diacetone alcohol)
40

Acid generators: PAG1 to PAG 6 of the following structural formulae

$$F_{2}C-SO_{3}^{-}$$

$$CF_{3}$$

$$PAG 4$$

$$I \longrightarrow CF_{2}$$

$$I \longrightarrow CF_{2}$$

$$CF_{3}$$

PAG 5

$$F_2C - SO_3^ CF_3$$

PAG 6

$$F_2C$$
— SO_3 -

-continued

Additive Quenchers 1 to 4:

Additive Quencher 2

25

40

Additive Quencher 1

Additive Quencher 3

Additive Quencher 4

Comparative Quenchers 1 to 6:

Comparative Quencher 2

Comparative Quencher 3

Comparative Quencher 4

Comparative Quencher 5

$$\begin{array}{c|c} & NH_2 & NH_2 \\ \hline & N^{\dagger}H_2 & \\ \hline & N^{\dagger}H_2 & \\ \hline & O \end{array}$$

Comparative Quencher 6

$$\begin{array}{c|c} & \text{NH}_2 & \text{NH}_2 \\ & \text{N} & \text{N}^+\text{H}_2 \end{array}$$

EUV Lithography Test

Each of the resist compositions in Tables 1 to 3 was spin coated on a silicon substrate having a 20-nm coating of silicon-containing spin-on hard mask SHB-A940 (Shin-Etsu Chemical Co., Ltd., silicon content 43 wt %) and prebaked on a hotplate at 105° C. for 60 seconds to form a resist film of 50 nm thick. Using an EUV scanner NXE3300 (ASML, NA 0.33, σ 0.9/0.6, quadrupole illumination), the resist film was exposed to EUV through a mask bearing a hole pattern at a pitch 46 nm (on-wafer size) and +20% bias. The resist film was baked (PEB) on a hotplate at the temperature
shown in Tables 1 to 3 for 60 seconds and developed in a 2.38 wt % TMAH aqueous solution for 30 seconds to form a hole pattern having a size of 23 nm in Examples 1 to 26

and Comparative Examples 1 to 6 or a dot pattern having a size of 23 nm in Example 27 and Comparative Example 7.

The resist pattern was observed under CD-SEM (CG-

The resist pattern was observed under CD-SEM (CG-5000, Hitachi High-Technologies Corp.). The exposure dose that provides a hole or dot pattern having a size of 23 nm is

reported as sensitivity. The size of 50 holes or dots at that dose was measured, from which a size variation (3σ) was computed and reported as CDU.

194

The resist composition is shown in Tables 1 to 3 together with the sensitivity and CDU of EUV lithography.

TABLE 1

Example	Polymer (pbw)	Acid generator (pbw)	Quencher (pbw)	Organic solvent (pbw)	PEB temp.	Sensitivity (mJ/cm ²)	CDU (nm)
1	Polymer 1 (100)	PAG 1 (26.2)	Quencher 1 (3.99)	PGMEA (2,000) DAA (500)	90	30	4.3
2	Polymer 1 (100)	PAG 2 (27.1)	Quencher 2 (4.69)	PGMEA (2,000) DAA (500)	90	28	4.0
3	Polymer 1 (100)	PAG 3 (25.4)	Quencher 3 (5.21)	PGMEA (2,000) DAA (500)	90	27	4.2
4	Polymer 1 (100)	PAG 4 (28.8)	Quencher 4 (4.55)	PGMEA (2,000) DAA (500)	90	26	4.3
5	Polymer 1 (100)	PAG 5 (26.5)	Quencher 5 (5.83)	PGMEA (2,000) DAA (500)	90	25	4.1
6	Polymer 1 (100)	PAG 6 (27.7)	Quencher 6 (3.85)	PGMEA (2.000) DAA (500)	90	27	4.0
7	Polymer 2 (100)	_	Quencher 7 (3.58)	PGMEA (400) CyH (2,000) PGME (100)	85	32	3.2
8	Polymer 2 (100)	_	Quencher 8 (5.04)	PGMEA (400) CyH (2,000) PGME (100)	85	30	3.3
9	Polymer 2 (100)	_	Quencher 9 (5.81)	PGMEA (400) CyH (2,000) PGME (100)	85	30	3.1
10	Polymer 2 (100)	_	Quencher 10 (5.16)	PGMEA (400) CyH (2,000) PGME (100)	85	31	3.1
11	Polymer 2 (100)	_	Quencher 11 (5.07) Additive Quencher 1 (2.36)	PGMEA (400) CyH (2,000) PGME (100)	85	32	3.0
12	Polymer 2 (100)	_	Quencher 12 (2.57) Additive Quencher 2 (2.36)	PGMEA (400) CyH (2,000) PGME (100)	85	28	3.1
13	Polymer 2 (100)	_	Quencher 13 (2.75) Additive Quencher 2 (3.81)	PGMEA (400) CyH (2,000) PGME (100)	85	27	3.1
14	Polymer 2 (100)	_	Quencher 11 (5.07) Additive Quencher 4 (4.46)	PGMEA (400) CyH (2,000) PGME (100)	85	29	3.3
15	Polymer 2 (100)	_	Quencher 14 (5.97)	PGMEA (400) CyH (2,000) PGME (100)	85	32	3.2
16	Polymer 2 (100)	_	Quencher 15 (5.31)	PGMEA (400) CyH (2,000) PGME (100)	85	31	3.3
17	Polymer 2 (100)	_	Quencher 16 (4.31)	PGMEA (400) CyH (2,000) PGME (100)	85	31	3.2
18	Polymer 2 (100)	_	Quencher 17 (4.45)	PGMEA (400) CyH (2,000) PGME (100)	85	32	3.2
19	Polymer 2 (100)	_	Quencher 18 (4.87)	PGMEA (400) CyH (2,000) PGME (100)	85	30	3.1
20	Polymer 2 (100)	_	Quencher 19 (4.37)	PGMEA (400) CyH (2,000) PGME (100)	85	31	3.3

TABLE 2

Example	Polymer (pbw)	Acid generator (pbw)	Quencher (pbw)	Organic solvent (pbw)	PEB temp. (° C.)	Sensitivity (mJ/cm ²)	CDU (nm)
21	Polymer 2 (100)	_	Quencher 20 (6.28)	PGMEA(400) CyH (2,000) PGME (100)	85	29	3.2
22	Polymer 2 (100)	_	Quencher 21 (5.41)	PGMEA (400) CyH (2,000) PGME (100)	85	29	3.1
23	Polymer 2 (100)	_	Quencher 22 (5.55)	PGMEA (400) CyH (2,000) PGME (100)	85	27	3.1
24	Polymer 2 (100)	_	Quencher 23 (5.67)	PGMEA (400) CyH (2,000) PGME (100)	85	28	3.0
25	Polymer 2 (100)	_	Quencher 24 (5.89)	PGMEA (400) DAA (2,000)	85	29	3.2
26	Polymer 3 (100)	_	Quencher 7 (3.58)	PGMEA (400) CyH (2,000) PGME (100)	80	27	3.3
27	Polymer 4 (100)	PAG 1 (12)	Quencher 7 (3.58)	PGMEA (2,000) DAA (500)	120	39	4.8

TABLE 3

Comparative Example	Polymer (pbw)	Acid generator (pbw)	Quencher (pbw)	Organic solvent (pbw)	PEB temp.	Sensitivity (mJ/cm ²)	CDU (nm)
1	Polymer 2	_	Comparative	PGMEA (400)	80	42	5.7
	(100)	_	Quencher 1 (1.91)	CyH (2,000) PGME (100)			
2	Polymer 2	_	Comparative	PGMEA (400)	80	43	5.9
	(100)		Quencher 2	CyH (2,000)			
			(3.13)	PGME (100)			
3	Polymer 2	_	Comparative	PGMEA (400)	80	42	4.9
	(100)		Quencher 3	CyH (2,000)			
			(3.64)	PGME (100)			
4	Polymer 2	_	Comparative	PGMEA (400)	80	40	4.4
	(100)		Quencher 4	CyH (2,000)			
			(3.43)	PGME (100)			
5	Polymer 2	_	Comparative	PGMEA (400)	80	42	5.3
	(100)		Quencher 5	CyH (2,000)			
			(2.93)	PGME (100)			
6	Polymer 2	_	Comparative	PGMEA (400)	80	42	5.2
	(100)		Quencher 6	CyH (2,000)			
			(3.41)	PGME (100)			
7	Polymer 4	PAG 4	Comparative	PGMEA (2,000)	120	52	6.5
	(100)	(12)	Quencher 6 (2.05)	DAA (500)			

It is demonstrated in Tables 1 to 3 that resist compositions 50 [3.3.3]undecane, biguanide or phosphazene compound comprising an iodized or brominated hydrocarbyl-containing carboxylic salt form patterns having a high sensitivity and reduced values of CDU.

Japanese Patent Application No. 2019-160863 is incorporated herein by reference.

Although some preferred embodiments have been described, many modifications and variations may be made thereto in light of the above teachings. It is therefore to be understood that the invention may be practiced otherwise than as specifically described without departing from the 60 wherein m and n are each independently 1, 2 or 3, scope of the appended claims.

The invention claimed is:

1. A resist composition comprising a base polymer and a salt, the salt consisting of an anion derived from a carboxylic acid having an iodized or brominated hydrocarbyl group 65 exclusive of iodized or brominated aromatic ring and a cation derived from a 2,5,8,9-tetraaza-1-phosphabicyclo

wherein the salt has the formula (A):

$$\left[\left(X^{\text{Bl}}\right)_{n}X^{1}\right]_{m}X^{2} \xrightarrow{O^{-}A^{+}}$$
(A)

 X^{B1} is iodine or bromine,

X¹ is a single bond, ether bond, ester bond, amide bond, carbonyl group or carbonate group,

 X^{2} is a single bond or a $\mathrm{C}_{1}\text{-}\mathrm{C}_{20}$ (m+1)-valent hydrocarbon group which may contain a heteroatom other than iodine and bromine,

R¹ is a C₁-C₂₀ (n+1)-valent aliphatic hydrocarbon group which may contain at least one moiety selected from the group consisting of fluorine, chlorine, hydroxyl, carboxyl, C₆-C₁₂ aryl, ether bond, ester bond, carbonyl, amide bond, carbonate, urethane bond, and urea bond,

A⁺ is a cation having the formula (A)-1, (A)-2 or (A)-3:

(A)-1 1(

(A)-2

20

30

(A)-3-1 ⁶⁰

 $\begin{array}{c|c}
R^{16} & & & & R^{19} \\
\hline
 & & & & & & \\
R^{15} & & & & & \\
\hline
 & & & & & & \\
N & & & & & & \\
\hline
 & & & & & & \\
N & & & & & & \\
\hline
 & & & & & & \\
R^{19} & & & & & \\
R^{2} & & & & & \\
\end{array}$

R²⁹ N P²⁴ R²⁴ R²⁵ R²⁵ N R²⁵ R²⁵

wherein R^{11} to R^{13} are each independently a C_1 - C_{24} hydrocarbyl group which may contain a heteroatom,

 R^{14} to R^{21} are each independently hydrogen or a C_1 - C_{24} hydrocarbyl group which may contain a heteroatom, a pair of R^{14} and R^{15} , R^{15} and R^{16} , R^{16} and R^{17} , R^{17} and R^{18} , R^{18} and R^{19} , R^{19} and R^{20} , or R^{20} and R^{21} may bond together to form a ring with the nitrogen atom to which they are attached, or the nitrogen atoms to which they are attached and intervening carbon atom, the ring optionally containing an ether bond,

 R^{22} to R^{29} are each independently hydrogen or a $C_1\text{-}C_{24}$ hydrocarbyl group which may contain a heteroatom, a pair of R^{22} and R^{23} , R^{23} and R^{24} , R^{24} and R^{25} , R^{25} and R^{26} , R^{26} and R^{27} , or R^{27} and R^{28} may bond together to 50 form a ring with the nitrogen atom to which they are attached, or the nitrogen atoms to which they are attached and intervening phosphorus atom, a pair of R^{22} and R^{23} , R^{24} and R^{25} , R^{26} and R^{27} , or R^{28} and R^{29} may bond together to form a group having the formula (A)-3-1, and R^{23} may be a group having the formula (A)-3-2 when R^{22} is hydrogen,

 R^{36} R^{37} R^{37} R^{39} R^{39} R^{39} R^{38}

wherein R³⁰ to R³⁹ are each independently hydrogen or a C₁-C₂₄ hydrocarbyl group which may contain a heteroatom, a pair of R³⁰ and R³¹, R³¹ and R³², R³² and R³³, R³³ and R³⁴, R³⁴ and R³⁵, R³⁶ and R³⁷, or R³⁸ and R³⁹ may bond together to form a ring with the nitrogen atom to which they are attached, or the nitrogen atoms to which they are attached and intervening phosphorus atom, a pair of R³⁰ and R³¹, R³² and R³³, or R³⁴ and R³⁵ may bond together to form a group having the formula (A)-3-1,

the broken line designates a valence bond.

2. The resist composition of claim 1, further comprising an acid generator capable of generating a sulfonic acid, sulfonimide or sulfonmethide.

3. The resist composition of claim 1, wherein the base polymer comprises recurring units having the formula (a1) or recurring units having the formula (a2):

$$V^2$$
 OR^{42}

wherein R⁴ is each independently hydrogen or methyl, R⁴¹ and R⁴² each are an acid labile group, Y¹ is a single bond, phenylene group, naphthylene group, or C₁-C₁₂ linking group containing at least one moiety selected from ester bond and lactone ring, and Y² is a single bond or ester bond.

4. The resist composition of claim **3** which is a chemically amplified positive resist composition.

5. The resist composition of claim 1 wherein the base polymer is free of an acid labile group.

6. The resist composition of claim **5** which is a chemically amplified negative resist composition.

7. The resist composition of claim 1 wherein the base polymer comprises recurring units of at least one type selected from recurring units having the formulae (f1) to (f3):

$$Z^{3}$$
 SO_{3}
 R^{56}
 SO_{3}
 R^{57}
 R^{58}

wherein R^A is each independently hydrogen or methyl, Z^1 is a single bond, phenylene group, $-O-Z^{11}$, -, $-C(=O)-O-Z^{11}$ or $-C(=O)-NH-Z^{11}$, Z^{11} is a C_1 - C_6 aliphatic hydrocarbylene group or phenylene group, which may contain a carbonyl, ester bond, ether bond or hydroxyl moiety,

 Z^2 is a single bond, $-Z^{21}$ —C(=O)—O—, $-Z^{21}$ —O— or $-Z^{21}$ —O—C(=O)—, Z^{21} is a C_1 - C_{12} saturated hydrocarbylene group which may contain a carbonyl moiety, ester bond or ether bond,

Z³ is a single bond, methylene, ethylene, phenylene, fluorinated phenylene, —C(=O)—O—Z³¹—, or —C(=O)—NH—Z³¹—, Z³¹ is a C¹-C₆ aliphatic hydrocarbylene group, phenylene group, fluorinated phenylene group, or trifluoromethyl-substituted phenylene group, which may contain a carbonyl moiety, ester bond, ether bond or hydroxyl moiety,

 R^{51} to R^{58} are each independently a C_1 - C_{20} hydrocarbyl group which may contain a heteroatom, any two of R^{53} , R^{54} and R^{55} or any two of R^{56} , R^{57} and R^{58} may bond together to form a ring with the sulfur atom to which they are attached,

R^{HF} is hydrogen or trifluoromethyl, and M⁻ is a non-nucleophilic counter ion.

- 8. The resist composition of claim 1, further comprising $_{20}$ an organic solvent.
 - 9. The resist composition of claim 1, further comprising a surfactant.
 - 10. The resist composition of claim 1 wherein R^1 is a C_1 - C_{20} (n+1)-valent straight or branched aliphatic hydrocarbon group which may contain at least one moiety selected from the group consisting of fluorine, chlorine, hydroxyl, carboxyl, C_6 - C_{12} aryl, ether bond, ester bond, carbonyl, amide bond, carbonate, urethane bond, and urea bond, when X^{BI} is bromine.
 - 11. A process for forming a pattern comprising the steps of applying the resist composition of claim 1 to form a resist film on a substrate, exposing the resist film to high-energy radiation, and developing the exposed resist film in a developer.
 - 12. The process of claim 11 wherein the high-energy radiation is ArF excimer laser radiation of wavelength 193 nm or KrF excimer laser radiation of wavelength 248 nm.
 - 13. The process of claim 11 wherein the high-energy radiation is EB or EUV of wavelength 3 to 15 nm.

* * * * *